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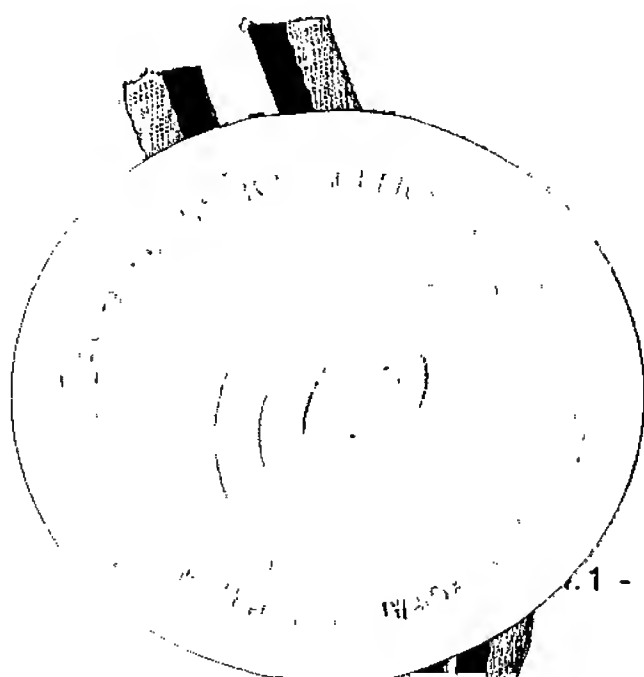
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Method and nucleic acids for the improved treatment of breast cell proliferative disorders

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Method and nucleic acids for the improved treatment of breast cell proliferative disorders

Field of the Invention

In American women, breast cancer is the most frequently diagnosed cancer and the second leading cause of cancer death. In women aged 40-55, breast cancer is the leading cause of death (Greenlee *et al.*, 2000). In 2002, there were 204,000 new cases of breast cancer in the US (data from the American Society of Clinical Oncology) and a comparable number in Europe.

Breast cancer is defined as the uncontrolled proliferation of cells within breasts tissues. Breasts are comprised of 15 to 20 lobes joined together by ducts. Cancer arises most commonly in the duct, but is also found in the lobes with the rarest type of cancer termed inflammatory breast cancer. It will be appreciated by those skilled in the art that there exists a continuing need to improve methods of early detection, classification and treatment of breast cancers. In contrast to the detection of some other common cancers such as cervical and dermal there are inherent difficulties in classifying and detecting breast cancers.

Due to current screening programs and the accessibility of this cancer to self-examination, breast cancer is diagnosed comparatively early: in about 93% of all newly diagnosed cases, the cancer has not yet metastasized, and in 65% of cases, even the lymph nodes are not yet affected..

The first step of any treatment is the assessment of the patient's condition comparative to defined classifications of the disease. However the value of such a system is inherently dependant upon the quality of the classification. Breast cancers are staged according to their size, location and occurrence of metastasis. Methods of treatment include the use

of surgery, radiation therapy, chemotherapy and endocrine therapy, which are also used as adjuvant therapies to surgery. Although the vast majority of early cancers are operable, i.e. the tumor can be completely removed by surgery, about one third of the patients with lymph-node negative diseases and about 50-60% of patients with node-positive disease will develop metastases during follow-up.

Based on this observation, systemic adjuvant treatment has been introduced for both node-positive and node-negative breast cancers. Systemic adjuvant therapy is administered after surgical removal of the tumor, and has been shown to reduce the risk of recurrence significantly (Early Breast Cancer Trialists' Collaborative Group, 1998). Several types of adjuvant treatment are available: endocrine treatment (for hormone receptor positive tumors), different chemotherapy regimens, and novel agents like Herceptin.

The growth of the majority of breast cancers (appr. 70-80%) is dependent on the presence of estrogen. Therefore, one important target for adjuvant therapy is the removal of estrogen (e.g. by ovarian ablation) or the blocking of its actions on the tumor cells (e.g. Tamoxifen). Endocrine treatment is thought to be efficient only in tumors that express hormone receptors (the estrogen receptor, ER, and/or the progesterone receptor, PR). Currently, the vast majority of women with hormone receptor positive breast cancer receive some form of endocrine treatment, independent of their nodal status. The most frequently used drug is Tamoxifen. However, even in hormone receptor positive patients, not all patients benefit from endocrine treatment. Adjuvant endocrine therapy reduces mortality rates by 22% while response rates to endocrine treatment in the advanced setting are 50 to 60% (Jordan et al., 2002, Jordan et al., 1999, Osborne et al., 1998, European Breast Cancer Cooperative Group, 1998).

Since Tamoxifen has relatively few side effects, treatment may be justified even for patients with low likelihood of benefit. However, these patients may require additional, more aggressive adjuvant treatment. This is supported by the fact that, even in earliest and least aggressive tumors, such as node-negative, hormone receptor positive tumours, about 21 % of patients relapse within 10 years after initial diagnosis if

they receive Tamoxifen monotherapy as adjuvant treatment (Early Breast Cancer Trialists Collaborative Group. Lancet, 1998).

Several cytotoxic regimens have shown to be effective in reducing the risk of relapse in breast cancer (Mansour *et al.*, 1998). According to current treatment guidelines, most node-positive patients receive adjuvant chemotherapy both in the US and Europe, since the risk of relapse is considerable.

Nevertheless, not all patients do relapse, and there is a proportion of patients who would never have relapsed even without chemotherapy, but who nevertheless receive chemotherapy due to the currently used criteria. In hormone receptor positive patients, chemotherapy is usually given before endocrine treatment, whereas hormone receptor negative patients receive only chemotherapy.

The situation for node-negative patients is particularly complex. In the US, cytotoxic chemotherapy is recommended for node-negative patients, if the tumor is larger than 1 cm. In Europe, chemotherapy is considered for the node-negative cases if one or more risk factors such as tumor size larger than 2 cm, negative hormone receptor status, or tumor grading of three or age <35 is present. In general, there is a tendency to select premenopausal women for additional chemotherapy whereas for postmenopausal women, chemotherapy is often omitted. Compared to endocrine treatment, in particular Tamoxifen, chemotherapy is highly toxic, with short-term side effects such as nausea, vomiting, bone marrow depression, and long-term effects such as cardiotoxicity and an increased risk for secondary cancers.

It is currently not clear which breast cancer patients should be selected for more aggressive therapy, although clinicians agree that there is a need for a subset of patients. The difficulty of selecting the right patients for chemotherapy, and the lack of suitable criteria is also reflected by a recent study which showed that chemotherapy is used much less frequently than recommended, based on data from the New Mexico Tumor registry (Du *et al.*, 2003). This study provides substantial evidence that there is a need for better selection

of patients for chemotherapy or other, more aggressive forms of breast cancer therapy.

The levels of observation that have been studied by the methodological developments of recent years in molecular biology, are the genes themselves, the translation of these genes into RNA, and the resulting proteins. The question of which gene is switched on at which point in the course of the development of an individual, and how the activation and inhibition of specific genes in specific cells and tissues are controlled is correlatable to the degree and character of the methylation of the genes or of the genome. In this respect, pathogenic conditions may manifest themselves in a changed methylation pattern of individual genes or of the genome.

DNA methylation plays a role, for example, in the regulation of the transcription, in genetic imprinting, and in tumorigenesis. Therefore, the identification of 5-methylcytosine as a component of genetic information is of considerable interest. However, 5-methylcytosine positions cannot be identified by sequencing since 5-methylcytosine has the same base pairing behaviour as cytosine. Moreover, the epigenetic information carried by 5-methylcytosine is completely lost during PCR amplification.

The currently most frequently used method for analysing DNA for 5-methylcytosine is based upon the specific reaction of bisulfite with cytosine which, upon subsequent alkaline hydrolysis, is converted to uracil which corresponds to thymine in its base pairing behaviour. However, 5-methylcytosine remains unmodified under these conditions. Consequently, the original DNA is converted in such a manner that methylcytosine, which originally could not be distinguished from cytosine by its hybridisation behaviour, can now be detected as the only remaining cytosine using "normal" molecular biological techniques, for example, by amplification and hybridisation or sequencing. All of these techniques are based on base pairing which can now be fully exploited. In terms of sensitivity, the prior art is defined by a method which encloses the DNA to be analysed in an

agarose matrix, thus preventing the diffusion and renaturation of the DNA (bisulfite only reacts with single-stranded DNA), and which replaces all precipitation and purification steps with fast dialysis (Olek A, Oswald J, Walter J. A modified and improved method for bisulphite based cytosine methylation analysis. *Nucleic Acids Res.* 1996 Dec 15;24(24):5064-6). Using this method, it is possible to analyse individual cells, which illustrates the potential of the method. However, currently only individual regions of a length of up to approximately 3000 base pairs are analysed, a global analysis of cells for thousands of possible methylation events is not possible. However, this method cannot reliably analyse very small fragments from small sample quantities either. These are lost through the matrix in spite of the diffusion protection.

An overview of the further known methods of detecting 5-methylcytosine may be gathered from the following review article: Rein, T., DePamphilis, M. L., Zorbas, H., *Nucleic Acids Res.* 1998, 26, 2255.

To date, barring few exceptions (e.g., Zeschnigk M, Lich C, Buiting K, Doerfler W, Horsthemke B. A single-tube PCR test for the diagnosis of Angelman and Prader-Willi syndrome based on allelic methylation differences at the SNRPN locus. *Eur J Hum Genet.* 1997 Mar-Apr;5(2):94-8) the bisulfite technique is only used in research. Always, however, short, specific fragments of a known gene are amplified subsequent to a bisulfite treatment and either completely sequenced (Olek A, Walter J. The pre-implantation ontogeny of the H19 methylation imprint. *Nat Genet.* 1997 Nov;17(3):275-6) or individual cytosine positions are detected by a primer extension reaction (Gonzalvo ML, Jones PA. Rapid quantitation of methylation differences at specific sites using methylation-sensitive single nucleotide primer extension (Ms-SNuPE). *Nucleic Acids Res.* 1997 Jun 15;25(12):2529-31, WO 95/00669) or by enzymatic digestion (Xiong Z, Laird PW. COBRA: a sensitive and quantitative DNA methylation assay. *Nucleic Acids Res.* 1997 Jun 15;25(12):2532-4). In addition, detection by hybridisation has also been described (Olek et al., WO 99/28498).

Further publications dealing with the use of the bisulfite technique for methylation detection in individual genes are: Grigg G, Clark S. Sequencing 5-methylcytosine residues in genomic DNA. *Bioessays*. 1994 Jun;16(6):431-6, 431; Zeschnigk M, Schmitz B, Dittrich B, Buiting K, Horsthemke B, Doerfler W. Imprinted segments in the human genome: different DNA methylation patterns in the Prader-Willi/Angelman syndrome region as determined by the genomic sequencing method. *Hum Mol Genet*. 1997 Mar;6(3):387-95; Feil R, Charlton J, Bird AP, Walter J, Reik W. Methylation analysis on individual chromosomes: improved protocol for bisulphite genomic sequencing. *Nucleic Acids Res*. 1994 Feb 25;22(4):695-6; Martin V, Ribieras S, Song-Wang X, Rio MC, Dante R. Genomic sequencing indicates a correlation between DNA hypomethylation in the 5' region of the pS2 gene and its expression in human breast cancer cell lines. *Gene*. 1995 May 19;157(1-2):261-4; WO 97/46705, WO 95/15373, and WO 97/45560.

An overview of the Prior Art in oligomer array manufacturing can be gathered from a special edition of *Nature Genetics* (*Nature Genetics Supplement*, Volume 21, January 1999), published in January 1999, and from the literature cited therein.

Fluorescently labelled probes are often used for the scanning of immobilised DNA arrays. The simple attachment of Cy3 and Cy5 dyes to the 5'-OH of the specific probe are particularly suitable for fluorescence labels. The detection of the fluorescence of the hybridised probes may be carried out, for example via a confocal microscope. Cy3 and Cy5 dyes, besides many others, are commercially available.

Matrix Assisted Laser Desorption Ionisation Mass Spectrometry (MALDI-TOF) is a very efficient development for the analysis of biomolecules (Karas M, Hillenkamp F. Laser desorption ionisation of proteins with molecular masses exceeding 10,000 daltons. *Anal Chem*. 1988 Oct 15;60(20):2299-301). An analyte is embedded in a light-absorbing matrix. The matrix is evaporated by a short laser pulse thus transporting the analyte molecule into the vapour phase in an unfragmented

manner. The analyte is ionised by collisions with matrix molecules. An applied voltage accelerates the ions into a field-free flight tube. Due to their different masses, the ions are accelerated at different rates. Smaller ions reach the detector sooner than bigger ones.

MALDI-TOF spectrometry is excellently suited to the analysis of peptides and proteins. The analysis of nucleic acids is somewhat more difficult (Gut I G, Beck S. DNA and Matrix Assisted Laser Desorption Ionization Mass Spectrometry. Current Innovations and Future Trends. 1995, 1; 147-57). The sensitivity to nucleic acids is approximately 100 times worse than to peptides and decreases disproportionally with increasing fragment size. For nucleic acids having a multiply negatively charged backbone, the ionisation process via the matrix is considerably less efficient. In MALDI-TOF spectrometry, the selection of the matrix plays an eminently important role. For the desorption of peptides, several very efficient matrixes have been found which produce a very fine crystallisation. There are now several responsive matrixes for DNA, however, the difference in sensitivity has not been reduced. The difference in sensitivity can be reduced by chemically modifying the DNA in such a manner that it becomes more similar to a peptide. Phosphorothioate nucleic acids in which the usual phosphates of the backbone are substituted with thiophosphates can be converted into a charge-neutral DNA using simple alkylation chemistry (Gut IG, Beck S. A procedure for selective DNA alkylation and detection by mass spectrometry. Nucleic Acids Res. 1995 Apr 25; 23(8): 1367-73). The coupling of a charge tag to this modified DNA results in an increase in sensitivity to the same level as that found for peptides. A further advantage of charge tagging is the increased stability of the analysis against impurities which make the detection of unmodified substrates considerably more difficult.

Genomic DNA is obtained from DNA of cell, tissue or other test samples using standard methods. This standard methodology is found in references such as Sambrook, Fritsch and Maniatis, Molecular Cloning: A Laboratory Manual, CSH Press, 2nd

edition, 1989: Isolation of genomic DNA from mammalian cells, Protocol I, p. 9.16 - 9.19. Also the manuals of several DNA extraction kits such as the QIAamp DNA mini kit give a good guidance on how to isolate genomic DNA.

Currently several predictive markers are under evaluation. As up to now most patients have received Tamoxifen as endocrine treatment most of the markers have been shown to be associated with response or resistance to tamoxifen. However, it is generally assumed that there is a large overlap between responders to one or the other endocrine treatment. In fact, ER and PR expression are used to select patients for any endocrine treatment. Among the markers which have been associated with TAM response is bcl-2. High bcl-2 levels showed promising correlation to TAM therapy response in patients with metastatic disease and prolonged survival and added valuable information to an ER negative patient subgroup (J Clin Oncology, 1997, 15 5: 1916-1922; Endocrine, 2000, 13(1):1-10). There is conflicting evidence regarding the independent predictive value of c-erbB2 (Her2/neu) overexpression in patients with advanced breast cancer that require further evaluation and verification (British J of Cancer, 1999, 79 (7/8):1220-1226; J Natl Cancer Inst, 1998, 90 (21): 1601-1608).

Other predictive markers include SRC-1 (steroid receptor coactivator-1), CGA gene over expression, cell kinetics and S phase fraction assays (Breast Cancer Res and Treat, 1998, 48:87-92; Oncogene, 2001, 20:6955-6959). Recently, uPA (Urokinase-type plasminogen activator) and PAI-1 (Plasminogen activator inhibitor type 1) together showed to be useful to define a subgroup of patients who have worse prognosis and who would benefit from adjuvant systemic therapy (J Clinical Oncology, 2002, 20 n° 4). However, all of these markers need further evaluations in prospective trials as none of them is yet a validated marker of response.

A number of cancer-associated genes have been shown to be inactivated by hypermethylation of CpG islands during breast tumorigenesis. Decreased expression of the calcium binding protein S100A2 (Accession number NM_005978) has been associated with the development of breast cancers. Hypermethylation of the promoter region of this gene has been observed in neoplastic cells thus providing evidence that

S100A2 repression in tumour cells is mediated by site-specific methylation.

The SYK gene (Accession number NM_003177) encodes a protein tyrosine kinase, Syk (spleen tyrosine kinase), that is highly expressed in hematopoietic cells. Syk is expressed in normal breast ductal epithelial cells but not in a subset of invasive breast carcinoma. Also, the loss of Syk expression seems to be associated with malignant phenotypes such as increased motility and invasion. The loss of expression occurs at the transcriptional level, and, as indicated by Yuan Y, Mendez R, Sahin A and Dai JL (Hypermethylation leads to silencing of the SYK gene in human breast cancer. Cancer Res. 2001 Jul 15;61(14):5558-61.), as a result of DNA hypermethylation.

The TGF- β type 2 receptor (encoded by the TGFBR2 gene, NM_003242) plays a role in trans-membrane signalling pathways via a complex of serine/threonine kinases. Mutations in the gene have been detected in some primary tumours and in several types of tumour-derived cell lines, including breast (Lucke CD, Philpott A, Metcalfe JC, Thompson AM, Hughes-Davies L, Kemp PR, Hesketh R. 'Inhibiting mutations in the transforming growth factor beta type 2 receptor in recurrent human breast cancer.' Cancer Res. 2001 Jan 15;61(2):482-5.).

The genes COX7A2L and GRIN2D were both identified as novel estrogen responsive elements by Watanabe et. al. (Isolation of estrogen-responsive genes with a CpG island library. Molec. Cell. Biol. 18: 442-449, 1998.) using the CpG-GBS (genomic binding site) method. The gene COX7A2L (Accession number NM_004718) encodes a polypeptide 2-like cytochrome C oxidase subunit VIIA. Northern blot analysis detected an upregulation of COX7A2L after estrogen treatment of a breast cancer cell line. The gene GRIN2D (Accession number NM_000836) encodes the N-methyl-D-aspartate, ionotropic, subunit 2D glutamate receptor, a subunit of the NMDA receptor channels associated with neuronal signalling. Furthermore expression of the cDNA has been observed in an osteosarcoma cell line. The gene VTN (also known as Vitronectin Accession number NM_000638) encodes a 75-kD glycoprotein (also called serum spreading factor or

complement S-protein) that promotes attachment and spreading of animal cells in vitro, inhibits cytolysis by the complement C5b-9 complex, and modulates antithrombin III-thrombin action in blood coagulation. Furthermore expression of this gene has been linked to progression and invasiveness of cancer cells.

The gene SFN (also known as Stratifin) encodes a polypeptide of the 14-3-3 family, 14-3-3 sigma. The 14-3-3 family of proteins mediates signal transduction by binding to phosphoserine-containing proteins. Expression of the SFN gene is lost in breast carcinomas, this is likely due to hypermethylation during the early stages of neoplastic transformation (see Umbricht CB, Evron E, Gabrielson E, Ferguson A, Marks J, Sukumar S. Hypermethylation of 14-3-3 sigma (stratifin) is an early event in breast cancer. Oncogene. 2001 Jun 7; 20(26):3348-53).

The gene PSAT1 (Accession number NM_021154) is not to be confused with the gene popularly referred to as PxySA (Accession number NM_001648) which encodes prostate specific antigen and whose technically correct name is kallikrein 3 . The gene PSAT1 encodes the protein phosphoserine aminotransferase which is the second step-catalysing enzyme in the serine biosynthesis pathway. Changes in gene expression levels have been monitored by mRNA expression analysis and upregulation of the gene has been identified in colonic carcinoma in a study of 6 samples (Electrophoresis 2002 Jun;23(11):1667-76 mRNA differential display of gene expression in colonic carcinoma.Ojala P, Sundstrom J, Gronroos JM, Virtanen E, Talvinen K, Nevalainen TJ).

The gene stathmin (NM_005563) codes for an oncoprotein 18, also known as stathmin, a conserved cytosolic phosphoprotein that regulates microtubule dynamics. The protein is highly expressed in a variety of human malignancies. In human breast cancers the stathmin gene has shown to be up-regulated in a subset of the tumours.

The gene PRKCD encodes a member of the family of protein kinase c enzymes, and is involved in B cell signaling and in

the regulation of growth, apoptosis, and differentiation of a variety of cell types.

Some of these molecules interact in a cascade-like manner. PRKCD activity that targets STMN1 is modulated by SFN binding and SYK phosphorylation. Together this influences tubulin polymerization that is required for cell division.

The gene MSMB (Accession number NM_002443) has been mapped to 10q11.2. It encodes the beta-microseminoprotein (MSP) which is one of the major proteins secreted by the prostate.

Furthermore, it may be useful as a diagnostic marker for prostate cancer. Using mRNA analysis low levels of beta-MSP mRNA expression and protein have been linked to progression under endocrine therapy and it has been postulated that it may be indicative of potentially aggressive prostate cancer (see Sakai H, Tsurusaki T, Kanda S, Koji T, Xuan JW, Saito Y 'Prognostic significance of beta-microseminoprotein mRNA expression in prostate cancer.' Prostate. 1999 Mar 1;38(4):278-84.).

The gene TP53 (Accession number NM_000546) encodes the protein p53, one of the most well characterised tumour suppressor proteins. The p53 protein acts as a transcription factor and serves as a key regulator of the cell cycle. Inactivation of this gene through mutation disrupts the cell cycle, which, in turn, assists in tumour formation. Methylation changes associated with this gene have been reported to be significant in breast cancer. Saraswati et. al. (Nature 405, 974 - 978 (22 Jun 2000) 'Compromised HOXA5 function can limit p53 expression in human breast tumours' reported that low levels of p53 mRNA in breast tumours was correlated to methylation of the HOXA5 gene. The product of the HOXA5 gene binds to the promoter region of the p53 and mediates expression of the gene. Methylation of the promoter region of the p53 gene itself has been reported (Kang JH, Kim SJ, Noh DY, Park IA, Choe KJ, Yoo OJ, Kang HS. 'Methylation in the p53 promoter is a supplementary route to breast carcinogenesis: correlation between CpG methylation in the p53 promoter and the mutation of the p53 gene in the progression from ductal carcinoma in

situ to invasive ductal carcinoma.' Lab Invest. 2001 Apr;81(4):573-9.). It was therein demonstrated that CpG methylation in the p53 promoter region is found in breast cancer and it was hypothesised that methylation in the p53 promoter region could be an alternative pathway to neoplastic progression in breast tumours. It has been observed that treatment with Tamoxifen decreases the level of expression of the p53 gene (Farczadi E, Kaszas I, Baki M, Szende B. 'Changes in apoptosis, mitosis, Her-2, p53 and Bcl2 expression in breast carcinomas after short-term tamoxifen treatment.' Neoplasma. 2002;49(2):101-3.)

The gene CYP2D6 (Accession number: NM_000106) is a member of the human cytochrome P450 (CYP) superfamily. Many members of this family are involved in drug metabolism (see for example Curr Drug Metab. 2002 Jun;3(3):289-309. Rodrigues AD, Rushmore TH.), of these Cytochrome P450 CYP2D6 is one of the most extensively characterised. It is highly polymorphic (more than 70 variations of the gene have been described), and allelic variation can result in both increased and decreased enzymatic activity. The CYP2D6 enzyme catalyses the metabolism of a large number of clinically important drugs including antidepressants, neuroleptics, some antiarrhythmics (Nature 1990 Oct 25;347(6295):773-6 Identification of the primary gene defect at the cytochrome P450 CYP2D locus. Gough AC, Miles JS, Spurr NK, Moss JE, Gaedigk A, Eichelbaum M, Wolf CR.).

The gene PTGS2 (Accession number NM_000963) encodes an inducible isozyme of prostaglandin-endoperoxide synthase (prostaglandin-endoperoxide synthase 2). It is also known as COX2 (cyclooxygenase 2). Aberrant methylation of this gene has been identified in lung carcinomas (Cancer Epidemiol Biomarkers Prev 2002 Mar;11(3):291-7 Hierarchical clustering of lung cancer cell lines using DNA methylation markers. Virmani AK, Tsou JA, Siegmund KD, Shen LY, Long TI, Laird PW, Gazdar AF, Laird-Offringa IA.).

The gene CGA (Accession number NM_000735) encodes the alpha polypeptide of glycoprotein hormones. Further, it has been identified as an estrogen receptor alpha (ER alpha)-responsive

gene and overexpression of the gene has been linked to ER positivity in breast tumours. Bieche *et. al.* examined mRNA levels of said gene in 125 ER alpha-positive postmenopausal breast cancer patients treated with primary surgery followed by adjuvant tamoxifen therapy. Initial results indicated significant links between CGA gene overexpression and Scarff-Bloom-Richardson histopathological grade I+II and progesterone and estrogen receptor positivity, which suggested that CGA is a marker of low tumour aggressiveness ('Identification of CGA as a Novel Estrogen Receptor-responsive Gene in Breast Cancer: An Outstanding Candidate Marker to predict the Response to Endocrine Therapy' *Cancer Research* 61, 1652-1658, February 15, 2001. Ivan Bièche, Béatrice Parfait, Vivianne Le Doussal, Martine Olivi, Marie-Christine Rio, Rosette Lidereau and Michel Vidaud). Further mRNA expression analysis linked CGA expression levels to Tamoxifen response, it was postulated that when combined with analysis of the marker ERBB2 (a marker of poor response) the gene may be useful as a predictive marker of tamoxifen responsiveness in breast cancer (*Oncogene* 2001 Oct 18;20(47):6955-9 The CGA gene as new predictor of the response to endocrine therapy in ER alpha-positive postmenopausal breast cancer patients. Bieche I, Parfait B, Nogues C, Andrieu C, Vidaud D, Spyrtos F, Lidereau R, Vidaud M.). The authors provided significant data associating the expression of the gene CGA with Tamoxifen treatment response. However, said analyses have all focused upon the analysis of relative levels of mRNA expression. This is not a methodology that is suitable for a medium or high throughput, nor is it a suitable basis for the development of a clinical assay.

The gene PITX2 (NM_000325) encodes the paired-like homeodomain transcription factor 2 which is known to be expressed during development of anterior structures such as the eye, teeth, and anterior pituitary. Although the expression of this gene is associated with cell differentiation and proliferation it has no heretofore recognised role in carcinogenesis or responsiveness to endocrine treatment. Toyota *et al.*, (2001. *Blood*. 97:2823-9.) found hypermethylation of the PITX2 gene in a large proportion of acute myeloid leukemias. Furthermore, this

hypermethylation is positively correlated to methylation of the ER gene.

RASSF1A (Ras association domain family 1A) gene is a candidate tumour suppressor gene at 3p21.3. The Ras GTPases are a superfamily of molecular switches that regulate cellular proliferation and apoptosis in response to extra-cellular signals. It is purported that *RASSF1A* is a tumour suppressor gene, and epigenetic alterations of this gene have been observed in a variety of cancers. Methylation of *RASSF1A* has been associated with poor prognosis in primary non-small cell lung cancer (Kim DH, Kim JS, Ji YI, Shim YM, Kim H, Han J, Park J., 'Hypermethylation of *RASSF1A* promoter is associated with the age at starting smoking and a poor prognosis in primary non-small cell lung cancer.' *Cancer Res.* 2003 Jul 1;63(13):3743-6.). It has also been associated with the development of pancreatic cancer (Kuzmin I, Liu L, Dammann R, Geil L, Stanbridge EJ, Wilczynski SP, Lerman MI, Pfeifer GP. 'Inactivation of RAS association domain family 1A gene in cervical carcinomas and the role of human papillomavirus infection.' *Cancer Res.* 2003 Apr 15;63(8):1888-93.), as well as testicular tumours and prostate carcinoma amongst others. The application of the methylation of this gene as a cancer diagnostic marker has been described in U.S. patent 6,596,488, it does not however describe its application in the selection of appropriate treatments regimens for patients.

Also located within 3p21 is the Dystroglycan precursor gene (Dystrophin-associated glycoprotein 1) (NM_004393). Dystroglycan (DG, also known as DAG1) is an adhesion molecule comprising two subunits namely alpha-DG and beta-DG. The molecule is responsible for crucial interactions between extracellular matrix and cytoplasmatic compartment and it has been hypothesised that as such it may contribute to progression to metastatic disease. Decreased expression of this gene has been associated with correlated with higher tumour grade and stage in colon, prostate and breast tumours.

The onecut-2 transcription factor gene (NM_004852) is located at 18q21.31 is a homeo-domain transcription factor regulator of liver gene expression in adults and during development.

The trefoil factor 1 (TFF1) gene (NM_003225) encodes a member of the trefoil family of proteins. The gene is also known as pS2. They are normally expressed at highest levels in the mucosa of the gastrointestinal tract, however they are often expressed ectopically in primary tumours of other tissues, including breast. The expression of TFF1 is regulated by estrogen in estrogen-responsive breast cancer cells in culture, its expression is associated with that of the estrogen receptor and TFF1 is a marker of hormone responsiveness in tumours (Schwartz et al., 1991. pS2 expression and response to hormonal therapy in patients with advanced breast cancer. Cancer Res. 51:624-8). TFF1 promoter methylation has been observed in nonexpressing gastric carcinoma-derived cell lines and tissues.

TMEFF2 (NM_016192) encodes a transmembrane protein containing an epidermal growth factor (EGF)-like motif and two follistatin domains. It has been shown to be overexpressed in prostate and brain tissues and it has been suggested that this is an androgen-regulated gene exhibiting antiproliferative effects in prostate cancer cells.

Methylation of the gene ESR1 (NM_000125), encoding the estrogen receptor has been linked to several cancer types including lung, oesophageal, brain and colorectal. The estrogen receptor (ESR) is a ligand-activated transcription factor composed of several domains important for hormone binding, DNA binding, and activation of transcription. Furthermore, it is the direct target of the anti-estrogenic compound Tamoxifen. Only tumours expressing estrogen receptor (ER+) can respond on Tamoxifen treatment.

The PCAF (NM_003884) gene encodes the p300/CBP-Associated Factor (PCAF). CBP and p300 are large nuclear proteins that bind to many sequence-specific factors involved in cell growth and/or differentiation. The p300/CBP associated factor

displays in vivo binding activity with CBP and p300. The protein has histone acetyl transferase activity with core histones and nucleosome core particles, indicating that it plays a direct role in transcriptional regulation. p300/CBP associated factor also associates with NF-kappa-B p65. This protein has been shown to regulate expression of the gene p53 by acetylation of Lys320 in the C-terminal portion of p53.

The WBP11 (NM_016312) gene encodes a nuclear protein, which co-localises with mRNA splicing factors and intermediate filament-containing perinuclear networks. It contains two proline-rich regions that bind to the WW domain of Npw38, a nuclear protein, and thus this protein is also called Npw38-binding protein NpwBP.

The TBC1 domain family, member 3 gene (TBC1D3, NM_032258) was discovered originally as an oncogene, also known as PRC17. The gene product contains a GTPase-activating protein (GAP) catalytic core motif and interacts directly with Rab5, stimulating its GTP hydrolysis. TBC1D3 is amplified in 15% of prostate cancers and highly overexpressed in approximately one-half of metastatic prostate tumors (Pei *et al.*, 2002; Cancer Res. 62:5420-4).

The CDK6 gene encodes a cyclin-dependent protein kinase regulating major cell cycle transitions in eukaryotic cells. The cdk6 kinase is associated with cyclins D1, D2, and D3 and can phosphorylate pRB, the product of the retinoblastoma tumor suppressor gene. The activation of cdk6 kinase occurs during mid-G1 (Meyerson and Harlow, 1994; Mol Cell Biol. 14:2077-86).

Description

In the following certain genetic regions are described for whom no genetic nomenclature is presently available. In each case the chromosomal location of the genetic sequence is denoted within parentetheses () and the genetic sequence is further described by its sequence according to Table 1.

The present invention provides methods and nucleic acids for the improved treatment planning of patients with cell

proliferative disorders of the breast tissues. The aim of the invention is achieved by assessment of one or both of two factors of particular relevance to patient treatment planning. The first factor is the characterisation of the cell proliferative disorder of the breast tissues and/or a metastases thereof in terms of aggressivity, the second factor being the prediction of disease free survival and/or response of a subject with said disorder to a therapy comprising one or more treatments which target the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion. Said treatments include, but are not limited to estrogen receptor modulators, estrogen receptor down-regulators, aromatase inhibitors, ovarian ablation, LHRH analogues and other centrally acting drugs influencing estrogen production. The prediction of response to a therapeutic regimen comprising one or more treatments which target the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion (a current treatment of choice as side effects are limited) further enables the physician to determine if additional treatments will be required in addition to or instead of this treatment. Treatments which may be used in addition to or instead of said treatment include, but are not limited to chemotherapy, radiotherapy, surgery, biological therapy, immunotherapy, antibodies and molecularly targeted drugs.

Characterisation of a breast cancer in terms of its predicted aggressiveness enables the physician to make an informed decision as to a therapeutic regimen with appropriate risk and benefit trade offs to the patient. Aggressiveness is taken to mean one or more of decreased patient survival or disease- or relapse-free survival, increased tumor-related complications and faster progression of tumor or metastases. According to the aggressiveness of the disease an appropriate treatment or treatments may be selected from the group consisting of chemotherapy, radiotherapy, surgery, biological therapy, immunotherapy, antibody treatments, treatments involving molecularly targeted drugs, estrogen receptor modulator treatments, estrogen receptor down-regulator treatments, aromatase inhibitors treatments, ovarian ablation, treatments providing LHRH analogues or other centrally acting drugs

influencing estrogen production. Wherein a cancer is characterised as 'aggressive' it is particularly preferred that a treatment such as, but not limited to, chemotherapy is provided in addition to or instead of an endocrine targeting therapy.

Using the methods and nucleic acids described herein, statistically significant models of patient disease free survival and/or responsiveness to treatment and/or disease progression can be developed and utilised to assist patients and clinicians in determining suitable treatment options to be included in the therapeutic regimen. In one aspect the described method is to be used to assess the utility of therapeutic regimens comprising one or more treatments which target the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion as a therapy for patients suffering from a cell proliferative disorder of the breast tissues. In particular this aspect of the method enables the physician to determine which treatments may be used in addition to or instead of said treatment. In a further aspect the described method enables the characterisation of the cell proliferative disorder in terms of aggressiveness, thereby enabling the physician to recommend suitable treatments. Thus, the present invention will be seen to reduce the problems associated with present breast cell proliferative disorder treatment response prediction methods.

Using the methods and nucleic acids as described herein, patient responsiveness can be evaluated before or during treatment for a cell proliferative disorder of the breast tissues, in order to provide critical information to the patient and clinician as to the likely progression of the disease. It will be appreciated, therefore, that the methods and nucleic acids exemplified herein can serve to improve a patient's quality of life and odds of treatment success by allowing both patient and clinician a more accurate assessment of the patient's treatment options.

The method according to the definition may be used for the improved treatment of all breast cell proliferative disorder patients, both pre and post menopausal and independant of

their node or estrogen receptor status. However, it is particularly preferred that said patients are node-negative and estrogen receptor positive.

The aim of the invention is most preferably achieved by means of the analysis of the methylation patterns of one or a combination of genes taken from the group taken from the group EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAU, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAU, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B and ESR1 (exon8) (see Table 1) and/or their regulatory regions.

The invention is characterised in that the nucleic acid of one or a combination of genes taken from the group EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAU, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAU, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B and ESR1 (exon8) are contacted with a reagent or series of reagents capable of distinguishing between methylated and non methylated CpG dinucleotides within the genomic sequence of interest.

The present invention makes available a method for the improved treatment and monitoring of breast cell proliferative disorders, by enabling the accurate prediction of a patient's disease free survival and/or response to treatment with a therapy comprising one or more treatments which target the estrogen receptor pathway or are involved in estrogen metabolism, production, or secretion.

In a particularly preferred embodiment, the method according to the invention enables the differentiation between patients who have a high probability of response to said therapy and

those who have a low probability of response to said therapy or a methylation characteristic predicted disease free survival time, in addition to the characterisation of tumors in terms of aggressiveness.

The method according to the invention may be used for the analysis of a wide variety of cell proliferative disorders of the breast tissues including, but not limited to, ductal carcinoma *in situ*, invasive ductal carcinoma, invasive lobular carcinoma, lobular carcinoma *in situ*, comedocarcinoma, inflammatory carcinoma, mucinous carcinoma, scirrhous carcinoma, colloid carcinoma, tubular carcinoma, medullary carcinoma, metaplastic carcinoma, and papillary carcinoma and papillary carcinoma *in situ*, undifferentiated or anaplastic carcinoma and Paget's disease of the breast.

The method according to the invention is particularly suited to the prediction of response to the aforementioned therapy in two treatment settings. In one embodiment, the method is applied to patients who receive endocrine pathway targeting treatment as secondary treatment to an initial non chemotherapeutical therapy, e.g. surgery (hereinafter referred to as the adjuvant setting) as illustrated in Figure 1. Such a treatment is often prescribed to patients suffering from Stage 1 to 3 breast carcinomas. In this embodiment patients disease free survival times are predicted according to their by detecting patients with worse disease free survival times the physician may choose to recommend the patient for further treatment, instead of or in addition to the endocrine targeting therapy(s), in particular but not limited to, chemotherapy.

In a further preferred embodiment said method is applied to patients suffering from a relapse of breast cancer following treatment by a primary means (preferably surgery) followed by a disease free period, and wherein the endocrine pathway targeting treatment has been prescribed in response to a detection of a relapse of the carcinoma. Such a treatment is often prescribed to patients suffering from later stage carcinomas, particularly wherein metastasis has occurred. Therefore this clinical setting shall also hereinafter be referred to as the 'metastatic setting'. In this embodiment

responders are those who enter partial or complete remission i.e. subjects whose cancer recedes to undetectable levels as opposed to those whose diseases further metastasise or remain above detectable levels. By detecting patients whose cancers are likely to metastasise the physician may choose to recommend the patient for further treatment, instead of or in addition to the endocrine targetting therapy(s), in particular but not limited to, chemotherapy.

This methodology presents further improvements over the state of the art in that the method may be applied to any subject, independent of the estrogen and/or progesterone receptor status. Therefore in a preferred embodiment, the subject is not required to have been tested for estrogen or progesterone receptor status.

The object of the invention is achieved by means of the analysis of the methylation patterns of one or more of the genes EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAU, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAU, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B, ESR1 (exon8) and/or their regulatory regions. In a particularly preferred embodiment the sequences of said genes comprise SEQ ID NOs: 1-61 and sequences complementary thereto.

The object of the invention may also be achieved by analysing the methylation patterns of one or more genes taken from the following subsets of said aforementioned group of genes. In one embodiment the object of the invention is the prediction of disease free survival and/or probability of response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion. This is achieved by analysis of the methylation patterns of one or more genes taken from the group consisting ERBB2, STMN1, TFF1, TMEFF2, ESR1, HSPB1, PITX2, COX7A2L, PLAU, VTN, PCAF, ONECUT2, BCL6, WBP11, TBC1D3, GRB7, CDK6, (Chr. 1p13.2), ABCA8 and

(Chr. 8q12.1) and wherein it is further preferred that the sequence of said genes comprise SEQ ID NOs: 5, 6, 12, 17, 18, 20, 23, 28, 16, 31, 33, 35, 36, 37, 43, 44, 46, 47, 49 and 51 respectively according to Table 1. In a further embodiment the object of the invention is the characterisation of the tumor in terms of aggressiveness. This is achieved by analysis of the methylation patterns of one or more genes taken from the group consisting APC, CSPG2, ERBB2, STK11, S100A2, TFF1, TGFBR2, TP53, TMEFF2, SYK, HSPB1, RASSF1, PSAT1, CGA, ESR2, ONECUT2, WBP11, CYP2D6, CDK6, ELK1, CGB and DAG1 , and wherein it is further preferred that the sequence of said genes comprise SEQ ID NOs: 2, 4, 5, 7, 11, 12, 13, 14, 17, 19, 20, 21, 25, 26, 29, 35, 37, 45, 46, 53, 55 and 59 respectively according to Table 1.

In a preferred embodiment said method is achieved by contacting said nucleic acid sequences in a biological sample obtained from a subject with at least one reagent or a series of reagents, wherein said reagent or series of reagents, distinguishes between methylated and non methylated CpG dinucleotides within the target nucleic acid.

In a preferred embodiment, the method comprises the following steps:

Preferably, said method comprises the following steps: In the *first step*, a sample of the tissue to be analysed is obtained. The source may be any suitable source, such as cell lines, histological slides, biopsies, tissue embedded in paraffin, bodily fluids, ejaculate, urine, blood and all possible combinations thereof. In a particularly preferred embodiment of the method said source is bodily fluids including post prostatic massage urine, ejaculate, urine, or blood. The DNA is then isolated from the sample. Extraction may be by means that are standard to one skilled in the art, including the use of commercially available kits, detergent lysates, sonification and vortexing with glass beads. Briefly, wherein the DNA of interest is encapsulated by a cellular membrane the biological sample must be disrupted and lysed by enzymatic, chemical or mechanical means. The DNA solution may then be cleared of proteins and other contaminants e.g. by digestion

with proteinase K. The genomic DNA is then recovered from the solution. This may be carried out by means of a variety of methods including salting out, organic extraction or binding of the DNA to a solid phase support. The choice of method will be affected by several factors including time, expense and required quantity of DNA.

Once the nucleic acids have been extracted, the genomic double stranded DNA is used in the analysis.

In the *second step* of the method, the genomic DNA sample is treated in such a manner that cytosine bases which are unmethylated at the 5'-position are converted to uracil, thymine, or another base which is dissimilar to cytosine in terms of hybridization behavior. This will be understood as 'pretreatment' herein.

The above-described treatment of genomic DNA is preferably carried out with bisulfite (hydrogen sulfite, disulfite) and subsequent alkaline hydrolysis that results in a conversion of non-methylated cytosine nucleobases to uracil or to another base that is dissimilar to cytosine in terms of base pairing behavior.

In the *third step* of the method, fragments of the pretreated DNA are amplified, using sets of primer oligonucleotides according to the present invention, and an amplification enzyme. The amplification of several DNA segments can be carried out simultaneously in one and the same reaction vessel. Typically, the amplification is carried out using a polymerase chain reaction (PCR). The set of primer oligonucleotides includes at least two oligonucleotides whose sequences are each reverse complementary, identical, or hybridize under stringent or highly stringent conditions to an at least 16-base-pair long segment of the base sequences of one or more of SEQ ID NO 206 to 449 and sequences complementary thereto.

In an alternate embodiment of the method, the methylation status of preselected CpG positions within the nucleic acid sequences comprising one or more of SEQ ID NO 1 to 61 may be detected by use of methylation-specific primer oligonucleotides. This technique (MSP) has been described in United States Patent No. 6,265,171 to Herman. The use of methylation status specific primers for the amplification of

bisulfite treated DNA allows the differentiation between methylated and unmethylated nucleic acids. MSP primers pairs contain at least one primer that hybridizes to a bisulfite treated CpG dinucleotide. Therefore, the sequence of said primers comprises at least one CpG dinucleotide. MSP primers specific for non-methylated DNA contain a "T" at the 3' position of the C position in the CpG. Preferably, therefore, the base sequence of said primers is required to comprise a sequence having a length of at least 9 nucleotides which hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 206-449 and sequences complementary thereto, wherein the base sequence of said oligomers comprises at least one CpG dinucleotide.

Wherein the method is for the prediction of probability of disease free survival and/or response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion it is particularly preferred that said nucleotide sequence(s) hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 70, 71, 192, 193, 72, 73, 194, 195, 84, 85, 206, 207, 94, 95, 216, 217, 96, 97, 218, 219, 100, 101, 222, 223, 106, 107, 228, 229, 116, 117, 238, 239, 92, 93, 214, 215, 122, 123, 244, 245, 126, 127, 248, 249, 130, 131, 252, 253, 132, 133, 254, 255, 134, 135, 256, 257, 146, 147, 268, 269, 148, 149, 270, 271, 152, 153, 274, 275, 154, 155, 276, 277, 158, 159, 280, 281, 162, 163, 284 and 285 said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

Wherein the method is for the the characterisation of the breast cell proliferative disorder in terms of aggressiveness it is particularly preferred that said nucleotide sequence(s) hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 64, 65, 186, 187, 68, 69, 190, 191, 70, 71, 192, 193, 74, 75, 196, 197, 82, 83, 204, 205, 84, 85, 206, 207, 86, 87, 208, 209, 88, 89, 210, 211, 94, 95, 216, 217, 98, 99, 220, 221, 100, 101, 222, 223, 102, 103, 224, 225, 110, 111, 232, 233, 112, 113, 234, 235, 118, 119, 240, 241, 130, 131, 252, 253, 134, 135, 256, 257, 150, 151, 272, 273, 152, 153, 274, 275, 166, 167, 288, 289, 170, 171, 292, 293, 178, 179, 300, 301, 148, 149, 270, 271, 150, 151, 272, 273, 152,

153, 274, 275, 154, 155, 276, 277, 156, 157, 278, 279, 158, 159, 280, 281, 160, 161, 282, 283, 162, 163, 284, 285, 164, 165, 286, 287, 166, 167, 288, 289, 168, 169, 290, 291, 170, 171, 292, 293, 172, 173, 294, 295, 174, 175, 296, 297, 176, 177, 298, 299, 178, 179, 300, 301, 180, 181, 302, 303, 182, 183, 304 and 305, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

A further preferred embodiment of the method comprises the use of *blocker* oligonucleotides. The use of such blocker oligonucleotides has been described by Yu et al., *BioTechniques* 23:714-720, 1997. Blocking probe oligonucleotides are hybridized to the bisulfite treated nucleic acid concurrently with the PCR primers. PCR amplification of the nucleic acid is terminated at the 5' position of the blocking probe, such that amplification of a nucleic acid is suppressed where the complementary sequence to the blocking probe is present. The probes may be designed to hybridize to the bisulfite treated nucleic acid in a methylation status specific manner. For example, for detection of methylated nucleic acids within a population of unmethylated nucleic acids, suppression of the amplification of nucleic acids which are unmethylated at the position in question would be carried out by the use of blocking probes comprising a 'CpA' or 'TpA' at the position in question, as opposed to a 'CpG' if the suppression of amplification of methylated nucleic acids is desired.

For PCR methods using blocker oligonucleotides, efficient disruption of polymerase-mediated amplification requires that blocker oligonucleotides not be elongated by the polymerase. Preferably, this is achieved through the use of blockers that are 3'-deoxyoligonucleotides, or oligonucleotides derivitized at the 3' position with other than a "free" hydroxyl group. For example, 3'-O-acetyl oligonucleotides are representative of a preferred class of blocker molecule.

Additionally, polymerase-mediated decomposition of the blocker oligonucleotides should be precluded. Preferably, such preclusion comprises either use of a polymerase lacking 5'-3' exonuclease activity, or use of modified blocker oligonucleotides having, for example, thioate bridges at the 5'-termini thereof that render the blocker molecule nuclease-

resistant. Particular applications may not require such 5' modifications of the blocker. For example, if the blocker- and primer-binding sites overlap, thereby precluding binding of the primer (e.g., with excess blocker), degradation of the blocker oligonucleotide will be substantially precluded. This is because the polymerase will not extend the primer toward, and through (in the 5'-3' direction) the blocker—a process that normally results in degradation of the hybridized blocker oligonucleotide.

A particularly preferred blocker/PCR embodiment, for purposes of the present invention and as implemented herein, comprises the use of peptide nucleic acid (PNA) oligomers as blocking oligonucleotides. Such PNA blocker oligomers are ideally suited, because they are neither decomposed nor extended by the polymerase.

Preferably, therefore, the base sequence of said *blocking oligonucleotides* is required to comprise a sequence having a length of at least 9 nucleotides which hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 206-449 , and sequences complementary thereto, wherein the base sequence of said oligonucleotides comprises at least one CpG, TpG or CpA dinucleotide.

Wherein the method is for the prediction of probability of disease free survival and/or response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion it is particularly preferred that said nucleotide sequence(s) hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 70, 71, 192, 193, 72, 73, 194, 195, 84, 85, 206, 207, 94, 95, 216, 217, 96, 97, 218, 219, 100, 101, 222, 223, 106, 107, 228, 229, 116, 117, 238, 239, 92, 93, 214, 215, 122, 123, 244, 245, 126, 127, 248, 249, 130, 131, 252, 253, 132, 133, 254, 255, 134, 135, 256, 257, 146, 147, 268, 269, 148, 149, 270, 271, 152, 153, 274, 275, 154, 155, 276, 277, 158, 159, 280, 281, 162, 163, 284 and 285, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

Wherein the method is for the the characterisation of the breast cell proliferative disorder in terms of aggressiveness it is particularly preferred that said nucleotide sequence(s)

hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 64, 65, 186, 187, 68, 69, 190, 191, 70, 71, 192, 193, 74, 75, 196, 197, 82, 83, 204, 205, 84, 85, 206, 207, 86, 87, 208, 209, 88, 89, 210, 211, 94, 95, 216, 217, 98, 99, 220, 221, 100, 101, 222, 223, 102, 103, 224, 225, 110, 111, 232, 233, 112, 113, 234, 235, 118, 119, 240, 241, 130, 131, 252, 253, 134, 135, 256, 257, 150, 151, 272, 273, 152, 153, 274, 275, 166, 167, 288, 289, 170, 171, 292, 293, 178, 179, 300, 301, 148, 149, 270, 271, 150, 151, 272, 273, 152, 153, 274, 275, 154, 155, 276, 277, 156, 157, 278, 279, 158, 159, 280, 281, 160, 161, 282, 283, 162, 163, 284, 285, 164, 165, 286, 287, 166, 167, 288, 289, 168, 169, 290, 291, 170, 171, 292, 293, 172, 173, 294, 295, 174, 175, 296, 297, 176, 177, 298, 299, 178, 179, 300, 301, 180, 181, 302, 303, 182, 183, 304 and 305, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

The fragments obtained by means of the amplification can carry a directly or indirectly detectable label. Preferred are labels in the form of fluorescence labels, radionuclides, or detachable molecule fragments having a typical mass that can be detected in a mass spectrometer. Where said labels are mass labels, it is preferred that the labeled amplificates have a single positive or negative net charge, allowing for better detectability in the mass spectrometer. The detection may be carried out and visualized by means of, e.g., matrix assisted laser desorption/ionization mass spectrometry (MALDI) or using electron spray mass spectrometry (ESI).

Matrix Assisted Laser Desorption/Ionization Mass Spectrometry (MALDI-TOF) is a very efficient development for the analysis of biomolecules (Karas & Hillenkamp, *Anal Chem.*, 60:2299-301, 1988). An analyte is embedded in a light-absorbing matrix. The matrix is evaporated by a short laser pulse thus transporting the analyte molecule into the vapour phase in an unfragmented manner. The analyte is ionized by collisions with matrix molecules. An applied voltage accelerates the ions into a field-free flight tube. Due to their different masses, the ions are accelerated at different rates. Smaller ions reach the detector sooner than bigger ones. MALDI-TOF spectrometry is well suited to the analysis of peptides and proteins. The analysis of nucleic acids is somewhat more

difficult (Gut & Beck, *Current Innovations and Future Trends*, 1:147-57, 1995). The sensitivity with respect to nucleic acid analysis is approximately 100-times less than for peptides, and decreases disproportionally with increasing fragment size. Moreover, for nucleic acids having a multiply negatively charged backbone, the ionization process via the matrix is considerably less efficient. In MALDI-TOF spectrometry, the selection of the matrix plays an eminently important role. For desorption of peptides, several very efficient matrixes have been found which produce a very fine crystallisation. There are now several responsive matrixes for DNA, however, the difference in sensitivity between peptides and nucleic acids has not been reduced. This difference in sensitivity can be reduced, however, by chemically modifying the DNA in such a manner that it becomes more similar to a peptide. For example, phosphorothioate nucleic acids, in which the usual phosphates of the backbone are substituted with thiophosphates, can be converted into a charge-neutral DNA using simple alkylation chemistry (Gut & Beck, *Nucleic Acids Res.* 23: 1367-73, 1995). The coupling of a charge tag to this modified DNA results in an increase in MALDI-TOF sensitivity to the same level as that found for peptides. A further advantage of charge tagging is the increased stability of the analysis against impurities, which makes the detection of unmodified substrates considerably more difficult.

In the *fourth step* of the method, the amplificates obtained during the third step of the method are analysed in order to ascertain the methylation status of the CpG dinucleotides prior to the treatment.

In embodiments where the amplificates were obtained by means of MSP amplification, the presence or absence of an amplificate is in itself indicative of the methylation state of the CpG positions covered by the primer, according to the base sequences of said primer.

Amplificates obtained by means of both standard and methylation specific PCR may be further analyzed by means of hybridization-based methods such as, but not limited to, array technology and probe based technologies as well as by means of techniques such as sequencing and template directed extension.

In one embodiment of the method, the amplicates synthesised in *step three* are subsequently hybridized to an array or a set of oligonucleotides and/or PNA probes. In this context, the hybridization takes place in the following manner: the set of probes used during the hybridization is preferably composed of at least 2 oligonucleotides or PNA-oligomers; in the process, the amplicates serve as probes which hybridize to oligonucleotides previously bonded to a solid phase; the non-hybridized fragments are subsequently removed; said oligonucleotides contain at least one base sequence having a length of at least 9 nucleotides which is reverse complementary or identical to a segment of the base sequences specified in the present Sequence Listing; and the segment comprises at least one CpG , TpG or CpA dinucleotide.

In a preferred embodiment, said dinucleotide is present in the central third of the oligomer. For example, wherein the oligomer comprises one CpG dinucleotide, said dinucleotide is preferably the fifth to ninth nucleotide from the 5'-end of a 13-mer. One oligonucleotide exists for the analysis of each CpG dinucleotide within the sequence according to SEQ ID NO 1 to 61, and the equivalent positions within SEQ ID NO 206-449 (according to Table 1). Said oligonucleotides may also be present in the form of peptide nucleic acids. The non-hybridized amplicates are then removed. The hybridized amplicates are then detected. In this context, it is preferred that labels attached to the amplicates are identifiable at each position of the solid phase at which an oligonucleotide sequence is located.

In yet a further embodiment of the method, the genomic methylation status of the CpG positions may be ascertained by means of oligonucleotide probes that are hybridised to the bisulfite treated DNA concurrently with the PCR amplification primers (wherein said primers may either be methylation specific or standard).

A particularly preferred embodiment of this method is the use of fluorescence-based Real Time Quantitative PCR (Heid et al., *Genome Res.* 6:986-994, 1996; also see United States Patent No. 6,331,393) employing a dual-labeled fluorescent oligonucleotide probe (TaqMan™ PCR, using an ABI Prism 7700 Sequence Detection System, Perkin Elmer Applied Biosystems,

Foster City, California). The TaqMan™ PCR reaction employs the use of a nonextendible interrogating oligonucleotide, called a TaqMan™ probe, which, in preferred embodiments, is designed to hybridize to a GpC-rich sequence located between the forward and reverse amplification primers. The TaqMan™ probe further comprises a fluorescent "reporter moiety" and a "quencher moiety" covalently bound to linker moieties (e.g., phosphoramidites) attached to the nucleotides of the TaqMan™ oligonucleotide. For analysis of methylation within nucleic acids subsequent to bisulfite treatment, it is required that the probe be methylation specific, as described in United States Patent No. 6,331,393, (hereby incorporated by reference in its entirety) also known as the MethyLight™ assay.

Variations on the TaqMan™ detection methodology that are also suitable for use with the described invention include the use of dual-probe technology (Lightcycler™) or fluorescent amplification primers (Sunrise™ technology). Both these techniques may be adapted in a manner suitable for use with bisulfite treated DNA, and moreover for methylation analysis within CpG dinucleotides.

A further suitable method for the use of probe oligonucleotides for the assessment of methylation by analysis of bisulfite treated nucleic acids. In a further preferred embodiment of the method, the *fifth step* of the method comprises the use of template-directed oligonucleotide extension, such as MS-SNuPE as described by Gonzalgo & Jones, *Nucleic Acids Res.* 25:2529-2531, 1997.

In yet a further embodiment of the method, the *fifth step* of the method comprises sequencing and subsequent sequence analysis of the amplificate generated in the *third step* of the method (Sanger F., et al., *Proc Natl Acad Sci USA* 74:5463-5467, 1977).

In one preferred embodiment of the method the nucleic acids according to SEQ ID NO 1 to 61, are isolated and treated according to the first three steps of the method outlined above, namely:

- a. obtaining, from a subject, a biological sample having subject genomic DNA;

- b. extracting or otherwise isolating the genomic DNA;
- c. treating the genomic DNA of b), or a fragment thereof, with one or more reagents to convert cytosine bases that are unmethylated in the 5-position thereof to uracil or to another base that is detectably dissimilar to cytosine in terms of hybridization properties;

and wherein the subsequent amplification of d) is carried out in a methylation specific manner, namely by use of methylation specific primers or *blocking oligonucleotides*, and further wherein the detection of the amplicates is carried out by means of a real-time detection probes, as described above. Wherein the subsequent amplification of d) is carried out by means of methylation specific primers, as described above, said methylation specific primers comprise a sequence having a length of at least 9 nucleotides which hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 206-449 , and sequences complementary thereto, wherein the base sequence of said oligomers comprises at least one CpG dinucleotide.

Wherein the method is for the prediction of disease free survival and/or probability of response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion it is particularly preferred that said *blocking oligonucleotide* nucleotide sequence(s) hybridizes to a pretreated nucleic acid sequence according to one of one of SEQ ID NO 70, 71, 192, 193, 72, 73, 194, 195, 84, 85, 206, 207, 94, 95, 216, 217, 96, 97, 218, 219, 100, 101, 222, 223, 106, 107, 228, 229, 116, 117, 238, 239, 92, 93, 214, 215, 122, 123, 244, 245, 126, 127, 248, 249, 130, 131, 252, 253, 132, 133, 254, 255, 134, 135, 256, 257, 146, 147, 268, 269, 148, 149, 270, 271, 152, 153, 274, 275, 154, 155, 276, 277, 158, 159, 280, 281, 162, 163, 284 and 285, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

Wherein the method is for the characterisation of the breast cell proliferative disorder in terms of aggressiveness it is particularly preferred that said *blocking oligonucleotide*

nucleotide sequence(s) hybridizes to a pretreated nucleic acid

sequence according to one of SEQ ID NO 64, 65, 186, 187, 68, 69, 190, 191, 70, 71, 192, 193, 74, 75, 196, 197, 82, 83, 204, 205, 84, 85, 206, 207, 86, 87, 208, 209, 88, 89, 210, 211, 94, 95, 216, 217, 98, 99, 220, 221, 100, 101, 222, 223, 102, 103, 224, 225, 110, 111, 232, 233, 112, 113, 234, 235, 118, 119, 240, 241, 130, 131, 252, 253, 134, 135, 256, 257, 150, 151, 272, 273, 152, 153, 274, 275, 166, 167, 288, 289, 170, 171, 292, 293, 178, 179, 300, 301, 148, 149, 270, 271, 150, 151, 272, 273, 152, 153, 274, 275, 154, 155, 276, 277, 156, 157, 278, 279, 158, 159, 280, 281, 160, 161, 282, 283, 162, 163, 284, 285, 164, 165, 286, 287, 166, 167, 288, 289, 168, 169, 290, 291, 170, 171, 292, 293, 172, 173, 294, 295, 174, 175, 296, 297, 176, 177, 298, 299, 178, 179, 300, 301, 180, 181, 302, 303, 182, 183, 304 and 305, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence. Step e) of the method, namely the detection of the specific amplicates indicative of the methylation status of one or more CpG positions according to SEQ ID NO 1 to 61 is carried out by means of real-time detection methods as described above.

In an alternative most preferred embodiment of the method the subsequent amplification of d) is carried out in the presence of *blocking oligonucleotides*, as described above. Said *blocking oligonucleotides* comprising a sequence having a length of at least 9 nucleotides which hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 206-449 and sequences complementary thereto, wherein the base sequence of said oligomers comprises at least one CpG, TpG or CpA dinucleotide. Step e) of the method, namely the detection of the specific amplicates indicative of the methylation status of one or more CpG positions according to SEQ ID NO 206-449 is carried out by means of real-time detection methods as described above.

In a further preferred embodiment of the method the nucleic acids according to SEQ ID NO 1 to 61 are isolated and treated according to the first three steps of the method outlined above, namely:

- a) obtaining, from a subject, a biological sample having subject genomic DNA;

- b) extracting or otherwise isolating the genomic DNA;
- c) treating the genomic DNA of b), or a fragment thereof, with one or more reagents to convert cytosine bases that are unmethylated in the 5-position thereof to uracil or to another base that is detectably dissimilar to cytosine in terms of hybridization properties; and wherein
- d) amplifying subsequent to treatment in c) is carried out in a methylation specific manner, namely by use of methylation specific primers or *blocking oligonucleotides*, and further wherein
- e) detecting of the amplicates is carried out by means of a real-time detection probes, as described above.

Wherein the subsequent amplification of c) is carried out by means of methylation specific primers, as described above, said methylation specific primers comprise a sequence having a length of at least 9 nucleotides which hybridizes to a pretreated nucleic acid sequence according to one of SEQ ID NO 206-449 and sequences complementary thereto, wherein the base sequence of said oligomers comprises at least one CpG dinucleotide. Wherein the method is for the prediction of disease free survival and/or probability of response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion it is particularly preferred that said methylation specific primers hybridize to a pretreated nucleic acid sequence according to one of one of SEQ ID NO 70, 71, 192, 193, 72, 73, 194, 195, 84, 85, 206, 207, 94, 95, 216, 217, 96, 97, 218, 219, 100, 101, 222, 223, 106, 107, 228, 229, 116, 117, 238, 239, 92, 93, 214, 215, 122, 123, 244, 245, 126, 127, 248, 249, 130, 131, 252, 253, 132, 133, 254, 255, 134, 135, 256, 257, 146, 147, 268, 269, 148, 149, 270, 271, 152, 153, 274, 275, 154, 155, 276, 277, 158, 159, 280, 281, 162, 163, 284 and 285, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

Wherein the method is for the characterisation of the breast cell proliferative disorder in terms of aggressiveness it is particularly preferred that said methylation specific primers

hybridize to a pretreated nucleic acid sequence according to one of SEQ ID NO 64, 65, 186, 187, 68, 69, 190, 191, 70, 71, 192, 193, 74, 75, 196, 197, 82, 83, 204, 205, 84, 85, 206, 207, 86, 87, 208, 209, 88, 89, 210, 211, 94, 95, 216, 217, 98, 99, 220, 221, 100, 101, 222, 223, 102, 103, 224, 225, 110, 111, 232, 233, 112, 113, 234, 235, 118, 119, 240, 241, 130, 131, 252, 253, 134, 135, 256, 257, 150, 151, 272, 273, 152, 153, 274, 275, 166, 167, 288, 289, 170, 171, 292, 293, 178, 179, 300, 301, 148, 149, 270, 271, 150, 151, 272, 273, 152, 153, 274, 275, 154, 155, 276, 277, 156, 157, 278, 279, 158, 159, 280, 281, 160, 161, 282, 283, 162, 163, 284, 285, 164, 165, 286, 287, 166, 167, 288, 289, 168, 169, 290, 291, 170, 171, 292, 293, 172, 173, 294, 295, 174, 175, 296, 297, 176, 177, 298, 299, 178, 179, 300, 301, 180, 181, 302, 303, 182, 183, 304 and 305, said contiguous nucleotides comprising at least one CpG, TpG or CpA dinucleotide sequence.

Additional embodiments of the invention provide a method for the analysis of the methylation status of genomic DNA according to the invention (SEQ ID NO 1 to 61) , and complements thereof) without the need for pretreatment.

Wherein the method is for the prediction of disease free survival and/or probability of response to a treatment which targets the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion it is particularly preferred that said genomic sequences are selected from SEQ ID NO 5, 6, 12, 17, 18, 20, 23, 28, 16, 31, 33, 35, 36, 37, 43, 44, 46, 47, 49 and 51.

Wherein the method is for the characterisation of the breast cell proliferative disorder in terms of aggressiveness it is particularly preferred that said genomic sequences are selected from SEQ ID NO 2, 4, 5, 7, 11, 12, 13, 14, 17, 19, 20, 21, 25, 26, 29, 35, 37, 45, 46, 53, 55 and 59.

In the *first step* of such additional embodiments, the genomic DNA sample is isolated from tissue or cellular sources.

Preferably, such sources include cell lines, histological slides, body fluids, or tissue embedded in paraffin. In the *second step*, the genomic DNA is extracted. Extraction may be by means that are standard to one skilled in the art, including but not limited to the use of detergent lysates, sonification and vortexing with glass beads. Once the nucleic

acids have been extracted, the genomic double-stranded DNA is used in the analysis.

In a preferred embodiment, the DNA may be cleaved prior to the treatment, and this may be by any means standard in the state of the art, in particular with methylation-sensitive restriction endonucleases.

In the *third step*, the DNA is then digested with one or more methylation sensitive restriction enzymes. The digestion is carried out such that hydrolysis of the DNA at the restriction site is informative of the methylation status of a specific CpG dinucleotide.

In the *fourth step*, which is optional but a preferred embodiment, the restriction fragments are amplified. This is preferably carried out using a polymerase chain reaction, and said amplicates may carry suitable detectable labels as discussed above, namely fluorophore labels, radionucleotides and mass labels.

In the *fifth step* the amplicates are detected. The detection may be by any means standard in the art, for example, but not limited to, gel electrophoresis analysis, hybridization analysis, incorporation of detectable tags within the PCR products, DNA array analysis, MALDI or ESI analysis.

When the methylation status of the selected CpG positions have been ascertained patient treatment relevant parameters can be ascertained wherein hypermethylation of the genes is associated with poor prognosis of said subject, aggressive characteristics of said cell proliferative disorder, poor disease free survival and/or lower probability of response of said subject to said treatment as relative to individuals with hypomethylation.

The term "hypermethylation" refers to the average methylation state corresponding to an *increased* (above average or median) presence of 5-mCyt at one or a plurality of CpG dinucleotides within a DNA sequence of a test DNA sample, relative to the amount of 5-mCyt found at corresponding CpG dinucleotides within a control DNA sample.

The term "hypomethylation" refers to the average methylation state corresponding to a *decreased* (below average

or median) presence of 5-mCyt at one or a plurality of CpG dinucleotides within a DNA sequence of a test DNA sample, relative to the amount of 5-mCyt found at corresponding CpG dinucleotides within a control DNA sample.

Kits

Moreover, an additional aspect of the present invention is a kit comprising, for example: a bisulfite-containing reagent; a set of primer oligonucleotides containing at least two oligonucleotides whose sequences in each case correspond, are complementary, or hybridize under stringent or highly stringent conditions to a 16-base long segment of the sequences SEQ ID NO: 1 to 61 and 206-449; oligonucleotides and/or PNA-oligomers; as well as instructions for carrying out and evaluating the described method. In a further preferred embodiment, said kit may further comprise standard reagents for performing a CpG position-specific methylation analysis, wherein said analysis comprises one or more of the following techniques: MS-SNuPE, MSP, MethyLight™, HeavyMethyl™, COBRA, and nucleic acid sequencing. However, a kit along the lines of the present invention can also contain only part of the aforementioned components.

Typical reagents (e.g., as might be found in a typical MethyLight®-based kit) for MethyLight® analysis may include, but are not limited to: PCR primers for specific gene (or methylation-altered DNA sequence or CpG island); TaqMan® probes; optimised PCR buffers and deoxynucleotides; and Taq polymerase.

Typical reagents (e.g., as might be found in a typical Ms-SNuPE-based kit) for Ms-SNuPE analysis may include, but are not limited to: PCR primers for specific gene (or methylation-altered DNA sequence or CpG island); optimised PCR buffers and deoxynucleotides; gel extraction kit; positive control primers; Ms-SNuPE primers for specific gene; reaction buffer (for the Ms-SNuPE reaction); and radioactive nucleotides. Additionally, bisulfite conversion reagents may include: DNA denaturation buffer; sulfonation buffer; DNA recovery reagents

or kit (e.g., precipitation, ultrafiltration, affinity column); desulfonation buffer; and DNA recovery components.

Typical reagents (e.g., as might be found in a typical MSP-based kit) for MSP analysis may include, but are not limited to: methylated and unmethylated PCR primers for specific gene (or methylation-altered DNA sequence or CpG island), optimized PCR buffers and deoxynucleotides, and specific probes.

In order to enable the disclosed method, the invention further provides the modified DNA of one or a combination of genes taken from the group EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAU, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAU, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B and ESR1 (exon8) as well as oligonucleotides and/or PNA-oligomers for detecting cytosine methylations within said genes. The present invention is based on the discovery that genetic and epigenetic parameters and, in particular, the cytosine methylation patterns of said genomic DNAs are particularly suitable for improved treatment and monitoring of breast cell proliferative disorders.

The nucleic acids according to the present invention can be used for the analysis of genetic and/or epigenetic parameters of genomic DNA.

This objective according to the present invention is achieved using a nucleic acid containing a sequence of at least 16 bases in length of the pretreated genomic DNA according to one of SEQ ID NO: 206 to SEQ ID NO: 449 and sequences complementary thereto.

The modified nucleic acids could heretofore not be connected with the improved treatment of breast cell proliferative disorders by prediction of disease free survival and/or

probability of response to treatment and/or characterisation of the disease in terms of aggressiveness.

The object of the present invention is further achieved by an oligonucleotide or oligomer for the analysis of pretreated DNA, for detecting the genomic cytosine methylation state, said oligonucleotide containing at least one base sequence having a length of at least 10 nucleotides which hybridises to a pretreated genomic DNA according to SEQ ID NO: 206 to SEQ ID NO: 449 . The oligomer probes according to the present invention constitute important and effective tools which, for the first time, make it possible to ascertain specific genetic and epigenetic parameters during the analysis of biological samples for features associated with a patient's disease free survival and/or response to endocrine treatment. Said oligonucleotides allow the improved treatment and monitoring of breast cell proliferative disorders. The base sequence of the oligomers preferably contains at least one CpG or TpG dinucleotide. The probes may also exist in the form of a PNA (peptide nucleic acid) which has particularly preferred pairing properties. Particularly preferred are oligonucleotides according to the present invention in which the cytosine of the CpG dinucleotide is within the middle third of said oligonucleotide e.g. the 5th - 9th nucleotide from the 5'-end of a 13-mer oligonucleotide; or in the case of PNA-oligomers, it is preferred for the cytosine of the CpG dinucleotide to be the 4th - 6th nucleotide from the 5'-end of the 9-mer.

The oligomers according to the present invention are normally used in so called "sets" which contain upto two oligomers and up to one oligomer for each of the CpG dinucleotides within SEQ ID NO: 206 to SEQ ID NO: 449 .

In the case of the sets of oligonucleotides according to the present invention, it is preferred that at least one oligonucleotide is bound to a solid phase. It is further preferred that all the oligonucleotides of one set are bound to a solid phase.

The present invention further relates to ~~a set of at least 2 n~~ (oligonucleotides and/or PNA-oligomers) used for detecting the cytosine methylation state of genomic DNA, by analysis of said sequence or treated versions of said sequence (of the genes EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAUI, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAUI, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B, ESR1 (exon8) as detailed in the sequence listing and Table 1) and sequences complementary thereto). These probes enable improved treatment and monitoring of breast cell proliferative disorders.

It will be obvious to one skilled in the art that the method according to the invention will be improved and supplemented by the incorporation of markers and clinical indicators known in the state of the art and currently used as predictive of the outcome of therapies which target endocrine or endocrine associated pathways. More preferably said markers include node status, age, menopausal status, grade, estrogen and progesterone receptors.

The genes that form the basis of the present invention may be used to form a "gene panel", i.e. a collection comprising the particular genetic sequences of the present invention and/or their respective informative methylation sites. The formation of gene panels allows for a quick and specific analysis of specific aspects of breast cancer treatment. The gene panel(s) as described and employed in this invention can be used with surprisingly high efficiency for the treatment of breast cell proliferative disorders by prediction of the outcome of treatment with a therapy comprising one or more drugs which target the estrogen receptor pathway or are involved in estrogen metabolism, production, or secretion. The analysis of each gene of the panel contributes to the evaluation of patient responsiveness, however, in a less preferred

embodiment the patient evaluation may be achieved by analysis of only a single gene. The analysis of a single member of the 'gene panel' would enable a cheap but less accurate means of evaluating patient responsiveness, the analysis of multiple members of the panel would provide a rather more expensive means of carrying out the method, but with a higher accuracy (the technically preferred solution).

The efficiency of the method according to the invention is improved when applied to patients who have not been treated with chemotherapy. Accordingly, it is a particularly preferred embodiment of the method wherein the method is used for the assessment of subjects who have not undergone chemotherapy.

According to the present invention, it is preferred that an arrangement of different oligonucleotides and/or PNA-oligomers (a so-called "array") made available by the present invention is present in a manner that it is likewise bound to a solid phase. This array of different oligonucleotide- and/or PNA-oligomer sequences can be characterised in that it is arranged on the solid phase in the form of a rectangular or hexagonal lattice. The solid phase surface is preferably composed of silicon, glass, polystyrene, aluminium, steel, iron, copper, nickel, silver, or gold. However, nitrocellulose as well as plastics such as nylon which can exist in the form of pellets or also as resin matrices are suitable alternatives.

Therefore, a further subject matter of the present invention is a method for manufacturing an array fixed to a carrier material for the improved treatment and monitoring of breast cell proliferative disorders. In said method at least one oligomer according to the present invention is coupled to a solid phase. Methods for manufacturing such arrays are known, for example, from US Patent 5,744,305 by means of solid-phase chemistry and photolabile protecting groups.

A further subject matter of the present invention relates to a DNA chip for the improved treatment and monitoring of breast cell proliferative disorders. The DNA chip contains at least

one nucleic acid according to the present invention. DNA chips are known, for example, in US Patent 5,837,832.

The oligomers according to the present invention or arrays thereof as well as a kit according to the present invention are intended to be used for the improved treatment and monitoring of breast cell proliferative disorders. According to the present invention, the method is preferably used for the analysis of important genetic and/or epigenetic parameters within genomic DNA, in particular for use in improved treatment and monitoring of breast cell proliferative disorders.

The methods according to the present invention are used, for improved treatment and monitoring of breast cell proliferative disorder by enabling more informed therapeutic regimens.

The present invention moreover relates to the diagnosis and/or prognosis of events which are disadvantageous or relevant to patients or individuals in which important genetic and/or epigenetic parameters within genomic DNA, said parameters obtained by means of the present invention may be compared to another set of genetic and/or epigenetic parameters, the differences serving as the basis for the diagnosis and/or prognosis of events which are disadvantageous or relevant to patients or individuals.

In the context of the present invention the term "hybridisation" is to be understood as a bond of an oligonucleotide to a completely complementary sequence along the lines of the Watson-Crick base pairings in the sample DNA, forming a duplex structure.

In the context of the present invention, "genetic parameters" are mutations and polymorphisms of genomic DNA and sequences further required for their regulation. To be designated as mutations are, in particular, insertions, deletions, point mutations, inversions and polymorphisms and, particularly preferred, SNPs (single nucleotide polymorphisms).

In the context of the present invention the term "methylation state" is taken to mean the degree of methylation present in a nucleic acid of interest, this may be expressed in absolute or relative terms i.e. as a percentage or other numerical value or by comparison to another tissue and therein described as hypermethylated, hypomethylated or as having significantly similar or identical methylation status.

In the context of the present invention the term "regulatory region" of a gene is taken to mean nucleotide sequences which affect the expression of a gene. Said regulatory regions may be located within, proximal or distal to said gene. Said regulatory regions include but are not limited to constitutive promoters, tissue-specific promoters, developmental-specific promoters, inducible promoters and the like. Promoter regulatory elements may also include certain enhancer sequence elements that control transcriptional or translational efficiency of the gene.

In the context of the present invention the term "chemotherapy" is taken to mean the use of drugs or chemical substances to treat cancer. This definition excludes radiation therapy (treatment with high energy rays or particles), hormone therapy (treatment with hormones or hormone analogues (synthetic substitutes) and surgical treatment.

In the context of the present invention, "epigenetic parameters" are, in particular, cytosine methylations and further modifications of DNA bases of genomic DNA and sequences further required for their regulation. Further epigenetic parameters include, for example, the acetylation of histones which, cannot be directly analysed using the described method but which, in turn, correlates with the DNA methylation.

In the context of the present invention the term "adjuvant treatment" is taken to mean a therapy of a cancer patient immediately following an initial non chemotherapeutical therapy, e.g. surgery. In general, the purpose of an adjuvant

therapy is to provide a significantly smaller risk of recurrences compared without the adjuvant therapy.

In the context of the present invention the term "estrogen and/or progesterone receptor positive" is taken to mean cells that express on their surface receptors that are susceptible to the binding of estrogens and/or progesterones.

While the present invention has been described with specificity in accordance with certain of its preferred embodiments, the following examples and figures serve only to illustrate the invention and is not intended to limit the invention within the principles and scope of the broadest interpretations and equivalent configurations thereof.

Figure 1 shows a preferred application of the method according to the invention. The X axis shows the tumour(s) mass, wherein the line '3' shows the limit of detectability. The Y-axis shows time. Accordingly said figure illustrates a simplified model of endocrine treatment of an Stage 1-3 breast tumour wherein primary treatment was surgery (at point 1), followed by adjuvant therapy with Tamoxifen. In a first scenario a responder to treatment (4) is shown as remaining below the limit of detectability for the duration of the observation. A non responder to the treatment (5) has a period of disease free survival (2) followed by relapse when the carcinoma mass reaches the level of detectability.

Figure 2 shows another preferred application of the method according to the invention. The X axis shows the tumour(s) mass, wherein the line '3' shows the limit of detectability. The Y-axis shows time. Accordingly said figure illustrates a simplified model of Endocrine treatment of an late stage breast tumour wherein primary treatment was surgery (at point 1), followed by relapse which is treated by Tamoxifen (2). In a first scenario a responder to treatment (4) is shown as remaining below the limit of detectability for the duration of the observation. A non responder to the treatment (5) does not recover from the relapse.

Figures 3 to 45 show the Kaplan-Meier estimated disease-free survival curves for single genes or oligonucleotide positions. The black plot shows the proportion of disease free patients in the population with above median methylation levels, the grey plot shows shows the proportion of disease free patients in the population with below median methylation levels

Figure 46 shows the methylation analysis of CpG islands according to Example 1. CpG islands per gene were grouped and their correlation with objective response determined by Hotelling's T^2 statistics. Black dots indicate the P -value of the indicated gene. The 20 most informative genes, ranked from left to right with increasing P -value, are shown. The top dotted line marks the uncorrected significance value ($P < 0.05$). The lower dotted line marks significance after false discovery rate correction of 25%. All genes with a P -value smaller or equal to the gene with the largest P -value that is below the lower line (in this case COX7A2L) are considered significant. The FDR correction chosen guarantees that the identified genes are with 75% chance true discoveries.

Figures 48 shows a ranked matrix of the best 11 amplificates of data obtained according to Example 1 (Metastatic setting, limited sample set). P -values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models. The figure is shown in greyscale, wherein the most significant CpG positions are at the bottom of the matrix with significance decreasing towards the top. Black indicates total methylation at a given CpG position, white represents no methylation at the particular position, with degrees of methylation represented in grey, from light (low proportion of methylation) to dark (high proportion of methylation). Each row represents one specific CpG position within a gene and each column shows the methylation profile for the different CpGs for one sample. The p -values for the individual CpG positions are shown on the right side. The p -values are the probabilities that the observed distribution occurred by chance in the data set.

Figure 49 shows a ranked matrix of some of the best markers obtained according to Example 1 (Metastatic setting, limited sample set). P-values were calculated from Likelihood ratio (LR) tests from univariate logistic regression models. The figure is shown in greyscale, wherein the most significant CpG positions are at the bottom of the matrix with significance decreasing towards the top. Black indicates total methylation at a given CpG position, white represents no methylation at the particular position, with degrees of methylation represented in grey, from light (low proportion of methylation) to dark (high proportion of methylation). Each row represents one specific CpG position within a gene and each column shows the methylation profile for the different CpGs for one sample. The p-values for the individual CpG positions are shown on the right side. The p-values are the probabilities that the observed distribution occurred by chance in the data set.

Figures 47 and 50 show the uncorrected p-values on a log-scale. P-values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models according to Example 1 (metastatic setting). Each individual genomic region of interest is represented as a point, the upper dotted line represents the cut off point for the 25% false discovery rate, the lower dotted line shows the Bonferroni corrected 5% limit.

Figure 51 shows a ranked matrix of the best 11 amplificates of data obtained according to Example 1 (Metastatic setting, all samples). P-values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models. The figure is shown in greyscale, wherein the most significant CpG positions are at the bottom of the matrix with significance decreasing towards the top. Black indicates total methylation at a given CpG position, white represents no methylation at the particular position, with degrees of methylation represented in grey, from light (low proportion of methylation) to dark (high proportion of methylation). Each

row represents one specific CpG position within a gene and each column shows the methylation profile for the different CpGs for one sample. The p-values for the individual CpG positions are shown on the right side. The p-values are the probabilities that the observed distribution occurred by chance in the data set.

Figure 52 shows the disease-free survival curves for a combination of two oligonucleotides each from the genes TBC1D3 and CDK6, and one oligonucleotide from the gene PITX2. The black plot shows the proportion of disease free patients in the population with above median methylation levels, the grey plot shows shows the proportion of disease free patients in the population with below median methylation levels

Figure 53 shows the plot according to Figure 52 and the classification of the sample set by means of the St. Gallen method. The unbroken lines represent the methylation analysis wherein the black plot shows the proportion of disease free patients in the population with above median methylation levels, the grey plot shows shows the proportion of disease free patients in the population with below median methylation levels. The broken lines represent the St. Gallen classification of the sample set wherein the black plot shows the disease free survival time of the high risk group and the grey plot shows the disease free survival of the low risk group.

Figure 54 shows the Kaplan-Meier estimated disease-free survival curves for a CpG position of the PITX2 gene by means of Real-Time methylation specific probe analysis. The lower plot shows the proportion of disease free patients in the population with above median methylation levels, the upper plot shows shows the proportion of disease free patients in the population with below median methylation levels. The X axis shows the disease free survival times of the patients in months, and the Y- axis shows the proportion of disease free survival patients.

SEQ ID NOS: 1 to 61 represent 5' and/or regulatory regions and/or CpG rich regions of the genes according to Table 1. These sequences are derived from Genbank and will be taken to include all minor variations of the sequence material which are currently unforeseen, for example, but not limited to, minor deletions and SNPs.

Example 1

DNA samples were extracted using the Wizzard Kit (Promega), samples from 278 patients were analysed, data analyses were carried out on a selection of candidate markers.

Bisulfite treatment and mPCR

Total genomic DNA of all samples was bisulfite treated converting unmethylated cytosines to uracil. Methylated cytosines remained conserved. Bisulfite treatment was performed with minor modifications according to the protocol described in Olek et al. (1996). After bisulfitation 10 ng of each DNA sample was used in subsequent mPCR reactions containing 6-8 primer pairs.

Each reaction contained the following:

2.5 pmol each primer
11.25 ng DNA (bisulfite treated)
Multiplex PCR Master mix (Qiagen)

Further details of the primers are shown in TABLE 2. Initial denaturation was carried out at 95°C for 15 min. Forty cycles were carried out as follows: Denaturation at 95°C for 30 sec, followed by annealing at 57°C for 90 sec., primer elongation at 72°C for 90 sec. A final elongation at 72°C was carried out for 10 min.

Hybridisation

All PCR products from each individual sample were then hybridised to glass slides carrying a pair of immobilised oligonucleotides for each CpG position under analysis. Each of these detection oligonucleotides was designed to hybridise to

the bisulphite converted sequence around one CpG site which was either originally unmethylated (TG) or methylated (CG). See Table 2 for further details of hybridisation oligonucleotides used. Hybridisation conditions were selected to allow the detection of the single nucleotide differences between the TG and CG variants.

5 μ l volume of each multiplex PCR product was diluted in 10 x Ssarc buffer. The reaction mixture was then hybridised to the detection oligonucleotides as follows. Denaturation at 95°C, cooling down to 10 °C, hybridisation at 42°C overnight followed by washing with 10 x Ssarc and dH2O at 42°C. Further details of the hybridisation oligonucleotides are shown in TABLE 3.

Fluorescent signals from each hybridised oligonucleotide were detected using genepix scanner and software. Ratios for the two signals (from the CG oligonucleotide and the TG oligonucleotide used to analyse each CpG position) were calculated based on comparison of intensity of the fluorescent signals.

Data analysis methods

Analysis of the chip data: From raw hybridisation intensities to methylation ratios; The log methylation ratio ($\log(\text{CG}/\text{TG})$) at each CpG position is determined according to a standardised preprocessing pipeline that includes the following steps: For each spot the median background pixel intensity is subtracted from the median foreground pixel intensity (this gives a good estimate of background corrected hybridisation intensities): For both CG and TG detection oligonucleotides of each CpG position the background corrected median of the 4 redundant spot intensities is taken; For each chip and each CpG position the $\log(\text{CG}/\text{TG})$ ratio is calculated; For each sample the median of $\log(\text{CG}/\text{TG})$ intensities over the redundant chip repetitions is taken. This ratio has the property that the hybridisation noise has approximately constant variance over the full range of possible methylation rates (Huber et al., 2002).

Hypothesis testing

The main task is to identify markers that show significant differences in the average degree of methylation between two classes. A significant difference is detected when the null hypothesis that the average methylation of the two classes is identical can be rejected with $p < 0.05$. Because we apply this test to a whole set of potential markers we have to correct the p-values for multiple testing. This was done by applying the False Discovery Rate (FDR) method (Dudoit et al., 2002).

For testing the null hypothesis that the methylation levels in the two classes are identical we used the likelihood ratio test for logistic regression models (Venables and Ripley, 2002). The logistic regression model for a single marker is a linear combination of methylation measurements from all CpG positions in the respective genomic region of interest (ROI). A significant p-value for a marker means that this ROI has some systematic correlation to the question of interest as given by the two classes. However, at least formally it makes no statement about the actual predictive power of the marker.

Logistic Regression

Logistic regression models are tools to model the probability of an event in dependence of one or more variables or factors. For example, if x denotes a specific methylation logratio, the probability that a patient responds to the applied therapy (Tamoxifen) is modeled as

$$P(\text{response} \mid x) = \exp(\beta_0 + \beta_1 x) / [1 + \exp(\beta_0 + \beta_1 x)]. \quad (1)$$

If x_1, \dots, x_k denote the k methylation logratios measured for one amplificate, the model is

$$P(\text{response} \mid x_1, \dots, x_k) = \exp(\beta_0 + \beta_1 x_1 + \dots + \beta_k x_k) / [1 + \exp(\beta_0 + \beta_1 x_1 + \dots + \beta_k x_k)]. \quad (2)$$

Significance of the respective amplificate is assessed using a likelihood-ratio test. This test calculates the difference of $-2 \log(\text{likelihood})$ for the full model and the null-model including just the intercept β_0 which is approximately χ^2

-distributed with k degrees of freedom under the null

hypotheses $\beta_1 = \dots = \beta_k = 0$.

If additional covariates are considered, the model contains an additional parameter for each covariate and the test statistic is calculated as the difference of $-2\text{Log}(\text{likelihood})$ or the full model and the null-model including intercept and covariates. Again, given the null hypothesis, this difference is approximately χ^2 -distributed with k degrees of freedom.

Ranked Matrices

For a graphical display of all group comparisons, ranked matrices are used. Each row represents one oligo pair, whereas each column of the matrix stands for one sample (or chip in the case of up- versus downmethyated Promega DNA comparisons). Oligo pairs are ranked according to their discriminatory power (Wilcoxon test, Fisher score or logistic regression), where the best "marker" is displayed on the bottom line.

Low methylation is displayed in light grey, high methylation in dark grey, and the data are normalized prior to display.

Cox Regression

Disease-free survival times (DFS) are modeled using Cox regression models. These models are similar to logistic regression models, but instead of probabilities, the hazard is modeled. The hazard gives the instantaneous risk of a relapse.

The models

$$h(t \mid x) = h_0(t) \cdot \exp(\beta x) \quad (3)$$

and

$$h(t \mid x_1, \dots, x_k) = h_0(t) \cdot \exp(\beta_1 x_1 + \dots + \beta_k x_k) \quad (4)$$

are used for uni- and multivariate analyses, respectively, where t is the time measured in months after surgery and $h_0(t)$ is the baseline hazard. Likelihood ratio tests are performed similar to those used for logistic regression. Again, the difference between

$2\text{Log}(\text{Likelihood})$ of full model and null-model is approximately χ^2

-distributed with k degrees of freedom

under the null hypotheses $\beta_1 = \dots = \beta_k = 0$.

Additional covariates can be included into the models.

Stepwise Regression Analysis

For both multivariate logistic and Cox regression models, a stepwise procedure is used in order to find submodels including only relevant variables. Two effects are usually achieved by these procedures: Variables (methylation ratios) that are basically unrelated to the dependent variable (response state or DFS, respectively) are excluded as they do not add relevant information to the model.

Out of a set of highly correlated variables, only the one with the the best relation to the dependent variable is retained. Inclusion of both types of variables can lead to numerical instabilities and a loss of power. Moreover, the predictive performance can be low due to overfitting. The applied algorithm aims at minimizing the Akaike information criterion (AIC) which is defined as

$$\text{AIC} = -2 \cdot \text{maximized log-likelihood} + 2 \cdot \text{\#parameters}.$$

The AIC is related to the predictive performance of a model, smaller values promise better performance.

Whereas the inclusion of additional variables always improves the model fit and thus increases the likelihood, the second term penalizes the estimation of additional parameters. The best model will present a compromise model with good fit and usually a small or moderate number of variables.

Results

Adjuvant setting

Analysis of the methylation patterns of patient samples treated with Tamoxifen as an adjuvant therapy immediately following surgery (see Figure 1) is shown in the plots according to Figures 3 to 45. For each amplificate, the mean methylation over all oligo-pairs for that amplificate was calculated and the population split into groups according to their mean methylation values, wherein one group was composed of individuals with a methylation score higher than the median and a second group composed of individuals with a methylation score lower than the median.

The results are shown in figures 3 to 9, as Cox model estimated disease-free survival curves. Figures 10 to 34 show

the disease free survival curves using the methylation analyses of only single oligonucleotide.

In a further analysis the recurrence of distant metastases only was analysed in figures 35 to 46.

The accuracy of the differentiation between the different groups was further increased by combining multiple oligonucleotides from different genes. Figures 52 show the combination of two oligonucleotides each from the genes TBC1D3 and CDK6, and one oligonucleotide from the gene PITX2. Figure 53 shows the classification of the patients from the sample set by means of the St. Gallen method (the current method of choice for estimating disease free survival) on top of Figure 52, thereby showing the improved effectiveness of methylation analysis over current methods, in particular post 80 months.

Metastatic setting

Analysis of the methylation patterns of patient samples treated with Tamoxifen in a metastatic setting (see Figure 2) is shown in the matrices according to Figures 46 to 52). The subjects analysed in this classification had relapsed following an initial treatment, the subsequent metastasis being treated by Tamoxifen.

In order to determine the ability of each gene promoter to predict success or failure of Tamoxifen treatment, the individual CpGs measured were combined per gene using Hotelling's T^2 statistics. Several genes were significantly associated with response to tamoxifen after correcting for multiple comparison with a moderate conservative false discovery rate of 25% (see Figure 52). The genes were ONECUT2, WBP11, CYP2D6, DAG1, ERBB2, S100A2, TFF1, TP53, TMEFF2, ESR1, SYK, RASSF1, PITX2, PSAT1, CGA and PCAF.

Figure 50 shows the uncorrected p-values on a log-scale. P-values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models. Each individual genomic region of interest is represented as a point, the upper dotted line represents the cut off point for the 25% false discovery rate, the lower dotted line shows the Bonferroni corrected 5% limit.

Figure 51 shows a ranked matrix of the best 11 amplificates of data obtained. P-values were calculated from Likelihood ratio

(LR) tests from multivariate logistic regression models. The figure is shown in greyscale, wherein the most significant CpG positions are at the bottom of the matrix with significance decreasing towards the top. Black indicates total methylation at a given CpG position, white represents no methylation at the particular position, with degrees of methylation represented in grey, from light (low proportion of methylation) to dark (high proportion of methylation). Each row represents one specific CpG position within a gene and each column shows the methylation profile for the different CpGs for one sample. The p-values for the individual CpG positions are shown on the right side. The p-values are the probabilities that the observed distribution occurred by chance in the data set.

Figures 47 through 49 the analysis of a subset of shows the uncorrected p-values on a log-scale.

Figure 47 shows the uncorrected p-values on a log-scale. P-values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models according to Example 1 (metastatic setting) . Each individual genomic region of interest is represented as a point, the upper dotted line represents the cut off point for the 25% false discovery rate, the lower dotted line shows the Bonferroni corrected 5% limit. Figure 48 shows a ranked matrix of the best 11 amplificates of data obtained. P-values were calculated from Likelihood ratio (LR) tests from multivariate logistic regression models. The figure is shown in greyscale, wherein the most significant CpG positions are at the bottom of the matrix with significance decreasing towards the top. Black indicates total methylation at a given CpG position, white represents no methylation at the particular position, with degrees of methylation represented in grey, from light (low proportion of methylation) to dark (high proportion of methylation). Each row represents one specific CpG position within a gene and each column shows the methylation profile for the different CpGs for one sample. The p-values for the individual CpG positions are shown on the right side. The p-values are the

probabilities that the observed distribution occurred by chance in the data set.

Real time Quantitative methylation analysis

Genomic DNA was analyzed using the Real Time PCR technique after bisulfite conversion. In this analysis four oligonucleotides were used in each reaction. Two non methylation specific PCR primers were used to amplify a segment of the treated genomic DNA containing a methylation variable oligonucleotide probe binding site. Two oligonucleotide probes competitively hybridise to the binding site, one specific for the methylated version of the binding site, the other specific to the unmethylated version of the binding site. Accordingly, one of the probes comprises a CpG at the methylation variable position (i.e. anneals to methylated bisulphite treated sites) and the other comprises a TpG at said position (i.e. anneals to unmethylated bisulphite treated sites). Each species of probe is labelled with a 5' fluorescent reporter dye and a 3' quencher dye wherein the CpG and TpG oligonucleotides are labelled with different dyes.

The reactions are calibrated by reference to DNA standards of known methylation levels in order to quantify the levels of methylation within the sample. The DNA standards were composed of bisulfite treated phi29 amplified genomic DNA (i.e. unmethylated), and/or phi29 amplified genomic DNA treated with Sss1 Methylase enzyme (thereby methylating each CpG position in the sample), which is then treated with bisulfite solution. Seven different reference standards were used with 0%, (i.e. phi29 amplified genomic DNA only), 5%, 10%, 25%, 50%, 75% and 100% (i.e. phi29 Sss1 treated genomic only).

The amount of sample DNA amplified is quantified by reference to the gene (β -actin (*ACTB*)) to normalize for input DNA. For standardization the primers and the probe for analysis of the *ACTB* gene lack CpG dinucleotides so that amplification is possible regardless of methylation levels. As there are no methylation variable positions, only one probe oligonucleotide is required.

The following oligonucleotides were used in the reaction:

Primer: TGGTGATGGAGGAGGTTTAGTAAGT (SEQ ID NO: 1088)

Primer: AACCAATAAAACCTACTCCTCCCTTAA (SEQ ID NO: 1089)

Probe: 6FAM-ACCACCACCCAACACACAATAACAAACACA-TAMRA or Dabcyl
(SEQ ID NO: 1090)

The extent of methylation at a specific locus was determined by the following formula:

$$\text{methylation rate} = 100 * I_{\text{CG}} / (I_{\text{CG}} + I_{\text{TG}})$$

(I = Intensity of the fluorescence of CG-probe or TG-probe)

Gene PITX2

Primers:

PITX2R02: GTAGGGGAGGGAAGTAGATGTT (SEQ ID NO: 1091)

PITX2Q02: TTCTAATCCTCCTTTCCACAATAA (SEQ ID NO: 1092)

Amplificate length : 143 bp

Probes:

PITX2cg1: FAM-AGTCGGAGTCGGGAGAGCGA-Darquencher (SEQ ID NO: 1093)

PITX2tg1: YAKIMA YELLOW-AGTTGGAGTTGGGAGAGTGAAAGGAGA-Darquencher (SEQ ID NO: 1094)

PCR components: 3 mM MgCl₂ buffer, 10x buffer, Hotstart TAQ

Program (45 cycles): 95 °C, 10 min

95 °C, 15 sec

62 °C, 1 min

Figure 54 shows the Kaplan-Meier estimated disease-free survival curves for a CpG position of the PITX2 gene by means of Real-Time methylation specific probe analysis. The lower plot shows the proportion of disease free patients in the population with above median methylation levels, the upper plot shows the proportion of disease free patients in the population with below median methylation levels. The x

axis shows the disease free survival times of the patients in months, and the Y- axis shows the proportion of disease free survival patients. The p-value (probability that the observed distribution occurred by chance) was calculated as 0.0031, thereby confirming the data obtained by means of array analysis according to figure 6.

Table 1

Accession no.	Gene name/locus	Genomic SEQ ID NO:	Pretreated methylated sequence (sense) SEQ ID NO:	Pretreated methylated strand (antisense) SEQ ID NO:	Pretreated unmethylated sequence (sense) SEQ ID NO:	Pretreated unmethylated sequence (antisense) SEQ ID NO:
NM_001965	EGR4	1	62	63	184	185
NM_000038	APC	2	64	65	186	187
NM_000077	CDKN2A	3	66	67	188	189
NM_004385	CSPG2	4	68	69	190	191
NM_004448	ERBB2	5	70	71	192	193
NM_005563	STMN1	6	72	73	194	195
NM_000455	STK11	7	74	75	196	197
NM_001216	CA9	8	76	77	198	199
NM_001604	PAX6	9	78	79	200	201
NM_006142	SFN	10	80	81	202	203
NM_005978	S100A2	11	82	83	204	205
NM_003225	TFF1	12	84	85	206	207
NM_003242	TGFBR2	13	86	87	208	209
NM_000546	TP53	14	88	89	210	211
NM_005427	TP73	15	90	91	212	213
NM_002658	PLAU	16	92	93	214	215
NM_016192	TMEFF2	17	94	95	216	217
NM_000125	ESR1	18	96	97	218	219
NM_003177	SYK	19	98	99	220	221
NM_001540	HSPB1	20	100	101	222	223
NM_007182	RASSF1	21	102	103	224	225
NM_015641	TES	22	104	105	226	227

NM_000325	PITX2	23	106	107	228	229
NM_000836	GRIN2D	24	108	109	230	231
NM_021154	PSAT1	25	110	111	232	233
NM_000735	CGA	26	112	113	234	235
NM_000106	CYP2D6	27	114	115	236	237
NM_004718	COX7A2L	28	116	117	238	239
NM_001437	ESR2	29	118	119	240	241
NM_002658	PLAU	30	120	121	242	243
NM_000638	VTN	31	122	123	244	245
NM_001055	SULT1A1	32	124	125	246	247
NM_003884	PCAF	33	126	127	248	249
NM_006254	PRKCD	34	128	129	250	251
NM_004852	ONECUT2	35	130	131	252	253
NM_001706	BCL6	36	132	133	254	255
NM_016312	WBP11	37	134	135	256	257
NM_002462	(MX1)	38	136	137	258	259
NM_138433	N.N.	39	138	139	260	261
NM_000484	APP	40	140	141	262	263
NM_002552	ORC4L	41	142	143	264	265
NM_138999	NETO1	42	144	145	266	267
NM_032258	TBC1D3	43	146	147	268	269
NM_005310	GRB7	44	148	149	270	271
NM_000106	CYP2D6	45	150	151	272	273
NM_001259	CDK6	46	152	153	274	275
	(Chr. 1p13.2)	47	154	155	276	277
	(Chr. 17q25.1)	48	156	157	278	279
NM_007168	ABCA8	49	158	159	280	281
	(Chr. 12q14.3)	50	160	161	282	283
	(Chr. 8q12.1)	51	162	163	284	285
NM_017490	MARK2	52	164	165	286	287
NM_005229	ELK1	53	166	167	288	289
	"Q8WUT3"	54	168	169	290	291

NM_000737	CGB	55	170	171	292	293
NM_001728	BSG	56	172	173	294	295
NM_005881	BCKDK	57	174	175	296	297
NM_014587	SOX8	58	176	177	298	299
NM_004393	DAG1	59	178	179	300	301
NM_020210	SEMA4B	60	180	181	302	303
NM_000125	ESR1 (exon8)	61	182	183	304	305

Table 2 Primers and amplicates according to Example 1

Gene:	Primer:	Amplicate Length:
EGR4 (SEQ ID NO: 1)	AGGGGGATTGAGTGTTAAGT (SEQ ID NO: 450) CCCAAACATAAACACAAAAT (SEQ ID NO: 451)	294
APC (SEQ ID NO: 2)	TCAACTACCATCAACTTCCTTA (SEQ ID NO: 452) AATTTATTTTGTAGTGTGTTAGTGGG (SEQ ID NO: 453)	491
CDKN2A (SEQ ID NO: 3)	GGGGTTGGTTGGTTATTAGA (SEQ ID NO: 454) AACCTCTACCCACCTAAAT (SEQ ID NO: 455)	256
CSPG2 (SEQ ID NO: 4)	GGATAGGAGTTGGGATTAAGAT (SEQ ID NO: 456) AAATCTTTTCAACACCAAAT (SEQ ID NO: 457)	414
ERBB2 (SEQ ID NO: 5)	GGAGGGGGTAGAGTTATTAGTT (SEQ ID NO: 458) TATACTTCCTCAAACAACCCTC (SEQ ID NO: 459)	257
STMN1 (SEQ ID NO: 6)	GAGTTTGTATTTAAGTTGAGTGGTT (SEQ ID NO: 460) AACAAAACAATACCCCTTCTAA (SEQ ID NO: 461)	334
STMN1	CCTCTTACTAACCTCAACCAAC	454

Gene:	Primer:	Amplificate Length:
(SEQ ID NO: 6)	(SEQ ID NO: 463) GAAAGGTAGGGAAGGATTTTT (SEQ ID NO: 462)	
STK11 (SEQ ID NO: 7)	TAAAAGAAGGATTTTTTGATTGG (SEQ ID NO: 464) CATCTTATTTACCTCCCTCCC (SEQ ID NO: 465)	528
CA9 (SEQ ID NO: 8)	GGGAAGTAGGTTAGGGTTAGTT (SEQ ID NO: 466) AAATCCTCCTCTCCAAATAAAT (SEQ ID NO: 467)	
PAX6 (SEQ ID NO: 9)	GGAGGGGAGAGGGTTATG (SEQ ID NO: 468) TACTATACACACCCCAAACAA (SEQ ID NO: 469)	374
SFN (SEQ ID NO: 10)	GAAGAGAGGAGAGGGAGGTA (SEQ ID NO: 470) CTATCCAACAAACCCAACA (SEQ ID NO: 471)	489
S100A2 (SEQ ID NO: 11)	GTTTTTAAGTTGGAGAAGAGGA (SEQ ID NO: 472) ACCTATAAATCACAACCCACTC (SEQ ID NO: 473)	460
TFF1 (SEQ ID NO: 12)	TTGGTGATGTTGATTAGAGTTT (SEQ ID NO: 474) TAAAACACCTTACATTTTCCCT (SEQ ID NO: 475)	449
TGFBR2 (SEQ ID NO: 13)	GTAATTTGAAGAAAGTTGAGGG (SEQ ID NO: 476) CCAACAATAAACAACCTCT (SEQ ID NO: 477)	296
TP53 (SEQ ID NO: 14)	TTGATGAGAAGAAAGGATTTAGT (SEQ ID NO: 478) TCAAATTCAATCAAAAACCTACC (SEQ ID NO: 479)	496
TP73 (SEQ ID NO: 15)	AGTAAATAGTGGGTGAGTTATGAA (SEQ ID NO: 480)	607

Gene:	Primer:	Amplificate Length:
	GAAAAACCTCTAAAACTACTCTCC (SEQ ID NO: 481)	
PLAU (SEQ ID NO: 16)	GAGAGAGATAGTTGGGGAGTTT (SEQ ID NO: 482) CAAACAACTTCATCTACCAAATAC (SEQ ID NO: 483)	453
TMEFF2 (SEQ ID NO: 17)	TGTTGGTTGTTGTTGTTGTT (SEQ ID NO: 484) CTTCTACCCATCCCAAAA (SEQ ID NO: 485)	319
ESR1 (SEQ ID NO: 18)	CTATCAATTCCCCCACTACT (SEQ ID NO: 487) TTGTTGGATAGAGGTTGAGTTT (SEQ ID NO: 486)	349
SYK (SEQ ID NO: 19)	GTGGGTTTTGGGTAGTTATAGA (SEQ ID NO: 488) TAACCTCCTCTCCTTACCAA (SEQ ID NO: 489)	485
HSPB1 (SEQ ID NO: 20)	CCTACCTCTACCACTTCTCAAT (SEQ ID NO: 491) AAGAGGGTTTAGTTTTTATTGG (SEQ ID NO: 490)	216
RASSF1 (SEQ ID NO: 21)	AGTGGGTAGGTTAAGTGTGTTG (SEQ ID NO: 492) CCCCAAAATCCAACTAAA (SEQ ID NO: 493)	319
TES (SEQ ID NO: 22)	AGGTTGGGGATTTTAGTTTTT (SEQ ID NO: 494) ACCTTCTTCACTTTATTTTCCA (SEQ ID NO: 495)	448
PITX2 (SEQ ID NO: 23)	TCCTCAACTCTACAAACCTAAAA (SEQ ID NO: 497) GTAGGGGAGGGAAGTAGATGT (SEQ ID NO: 496)	408
GRIN2D (SEQ ID NO: 24)	ATAGTTTGTGGTTTGGATTTTT (SEQ ID NO: 498) AAAACCTTCCCTAACTTCAAT	435

Gene:	Primer:	Amplificate Length:
	(SEQ ID NO: 499)	
PSAT1 (SEQ ID NO: 25)	GTAGGTGGTTAATTTTGGGTT (SEQ ID NO: 500) CTCATTCACACTATATCCATTCA (SEQ ID NO: 501)	500
PSAT1 (SEQ ID NO: 25)	TAAGAGAGAGGAGTTGAGGTTT (SEQ ID NO: 502) CCAAAATTAACCACCTACCTAA (SEQ ID NO: 503)	478
CGA (SEQ ID NO: 26)	TAGTGGTATAAGTTTGGAAATGTT (SEQ ID NO: 504) TCCACCTACATCTAAACCCTAA (SEQ ID NO: 505)	364
CYP2D6 (SEQ ID NO: 27)	CCTCCTAAACTAAATCCAACAA (SEQ ID NO: 507) GGGGTTAAGGTTTTTATGGTA (SEQ ID NO: 506)	418
COX7A2L (SEQ ID NO: 28)	AATCCTAAAAACCCTAACTTTTAAT (SEQ ID NO: 509) GGAGGTGTAAGGAGAATAGAGA (SEQ ID NO: 508)	398
ESR2 (SEQ ID NO: 29)	AAACCTTCCCAATAACCTCTTA (SEQ ID NO: 511) TAGAGGGGAGTAGTGTTTGAGT (SEQ ID NO: 510)	471
PLAU (SEQ ID NO: 30)	GTGATATTTGGGGATTGTTATT (SEQ ID NO: 512) ACTCCCTCCCCTATCTTACA (SEQ ID NO: 513)	479
VTN (SEQ ID NO: 31)	GTTATTTGGGTTAATGTAGGGA (SEQ ID NO: 514) TCTATCCCCTCAAACCTAAAAA (SEQ ID NO: 515)	492
SULT1A1 (SEQ ID NO: 32)	ATACTACCAAACCCACTCAAAC (SEQ ID NO: 517) GAATTAGGGAAGGAGTTAGTTG (SEQ ID NO: 516)	448

Gene:	Primer:	Amplificate Length:
PCAF (SEQ ID NO: 33)	GGATAAATGATTGAGAGGTTGT (SEQ ID NO: 518) CCTCCCTTAATTCTCCTACC (SEQ ID NO: 519)	369
PRKCD (SEQ ID NO: 34)	CTTAACCCATCCCAATCA (SEQ ID NO: 521) GATAGAAGGATTTTAGTTTATTGTT (SEQ ID NO: 520)	322
ONECUT2 (SEQ ID NO: 35)	TTTGTTGGGATTTGTTAGGAT (SEQ ID NO: 522) AAACATTTTACCCCTCTAAACC (SEQ ID NO: 523)	467
BCL6 (SEQ ID NO: 36)	CATCACCCTTCTAAAAACCC (SEQ ID NO: 525) GGGTAAGAAAGAAGGAATTAGTTT (SEQ ID NO: 524)	456
WBP11 (SEQ ID NO: 37)	AAGAGGTGAGGAAGAGTAGTAAAT (SEQ ID NO: 526) CTCCCAACAATAAATCAAAAT (SEQ ID NO: 527)	437
(MX1) (SEQ ID NO: 38)	TGTAGGAGAGGTTGGGAAG (SEQ ID NO: 528) CCAAACATAACATCCACTAAAA (SEQ ID NO: 529)	341
N.N. (SEQ ID NO: 39)	TAGGTTTAAGAGGAGAGGGAAT (SEQ ID NO: 530) AAACAACCTACCCAAATCCAAC (SEQ ID NO: 531)	433
APP (SEQ ID NO: 40)	GAGTAAGGAAGGGGGATG (SEQ ID NO: 532) AACCCAAATCTTTAATACAAAA (SEQ ID NO: 533)	494
NETO1 (SEQ ID NO: 42)	GGAGTTTTTAGAAGAGGAAGATT (SEQ ID NO: 534) ACTTCACAATAAATACCCTCCC (SEQ ID NO: 535)	395
TBC1D3	GGTAGAGGAAGTAGTTGGTTTG	490

Gene:	Primer:	Amplificate Length:
(SEQ ID NO: 43)	(SEQ ID NO: 536) CTTTTATATTTCTCCCAATCTCC (SEQ ID NO: 537)	
GRB7 (SEQ ID NO: 44)	AAAATCCATAACCACCAAATA (SEQ ID NO: 539) TTAGGAAGTTT TAGGAATGAGG (SEQ ID NO: 538)	416
CYP2D6 (SEQ ID NO: 45)	AATTCCTAACCCACTATCCTC (SEQ ID NO: 541) ATTTGTAGTTTGGGGTGATTT (SEQ ID NO: 540)	379
CDK6 (SEQ ID NO: 46)	ACCTTAAACACCTTCCCATAA (SEQ ID NO: 543) GTGTAATGATTTTGGATTGAGA (SEQ ID NO: 542)	456
(Chr. 1p13.2) (SEQ ID NO: 47)	AAGGAAGGTAGAGGGTTGAGT (SEQ ID NO: 544) AAAATCCAAAATTAACACCATT (SEQ ID NO: 545)	499
(Chr. 17q25.1) (SEQ ID NO: 48)	AGTAGATGAAGTTGGGGATTAG (SEQ ID NO: 546) TCCTACTATCCCTTCTCAAAAA (SEQ ID NO: 547)	500
ABCA8 (SEQ ID NO: 49)	TGATTGTGTAGATTATTTTGGTT (SEQ ID NO: 548) CAAACCTCTTAAACCTCAATCTC (SEQ ID NO: 549)	499
(Chr. 12q14.3) (SEQ ID NO: 50)	ACCCTAACATTCTCTAAACAACA (SEQ ID NO: 551) GATGAAAGTGGAAAGATTATGG (SEQ ID NO: 550)	441
(Chr. 8q12.1) (SEQ ID NO: 51)	CTCCAACCTCTCCTCACCTC (SEQ ID NO: 553) ATTTGAAGGTTGTGTTTGTAGA (SEQ ID NO: 552)	343
MARK2 (SEQ ID NO: 52)	TCACCACTATCCTCAATAATCA (SEQ ID NO: 555)	476

Gene:	Primer:	Amplificate Length:
	TAAAGTAGGAAGGTTTGGTTTG (SEQ ID NO: 554)	
ELK1 (SEQ ID NO: 53)	CCTCTAATTCCTATCAATCACC (SEQ ID NO: 557) TTAGAAGTGAAAGTAGAAGGGTTT (SEQ ID NO: 556)	435
Q8WUT3 (SEQ ID NO: 54)	GGTTAGAAGTTAGAGGGGTAGG (SEQ ID NO: 558) CCATCCCATTTACCTATAAAAAT (SEQ ID NO: 559)	406
CGB (SEQ ID NO: 55)	TCCACCCTATTTTCTACCAA (SEQ ID NO: 561) TTTGTTTTAGGTGGTGTGTAAT (SEQ ID NO: 560)	417
BSG (SEQ ID NO: 56)	TTATCTATCCCCACACCCTAAT (SEQ ID NO: 563) GGAGTAGGTGAGGAGTATTTTG (SEQ ID NO: 562)	420
BCKDK (SEQ ID NO: 57)	TCACCTCCTTTTACAACCAAT (SEQ ID NO: 565) TTTGGGAGAGTTTTAGGATTTA (SEQ ID NO: 564)	258
SOX8 (SEQ ID NO: 58)	GGGTGGGTAGTAGGTTTGTT (SEQ ID NO: 566) ACACACTCCTTAAACTCTTCC (SEQ ID NO: 567)	435
DAG1 (SEQ ID NO: 59)	AATACCAACCCAAACATCTACC (SEQ ID NO: 569) TTTGTTTATGTGGAGTTTATTGT (SEQ ID NO: 568)	315
ORC4L (SEQ ID NO: 41)	CACTCAAACTTCCCTACCTAC (SEQ ID NO: 571) GGTAATGGTGGGGGTAAAT (SEQ ID NO: 570)	489
SEMA4B (SEQ ID NO: 60)	ACCAAAATACTACTCCCAAATC (SEQ ID NO: 573) GGGTAGAGGGAGGTTATTGTT	337

Gene:	Primer:	Amplificate Length:
	(SEQ ID NO: 572)	
ESR1 (exon8) (SEQ ID NO: 61)	TATGATTTGTTGTTGGAGATGT (SEQ ID NO: 574) CTTAAAATCCCTTTAACTATTCCC (SEQ ID NO: 575)	388

Table 3 Hybridisation oligonucleotides according to Example 1

Gene	Oligo:
ONECUT2 (SEQ ID NO: 35)	TACGTAGTTGCGCGTT (SEQ ID NO: 800)
ONECUT2 (SEQ ID NO: 35)	GTATGTAGTTGTGTGTT (SEQ ID NO: 801)
ONECUT2 (SEQ ID NO: 35)	TTTTGTGCGTACGGAT (SEQ ID NO: 802)
ONECUT2 (SEQ ID NO: 35)	TTTTTGTGTGTATGGAT (SEQ ID NO: 803)
ONECUT2 (SEQ ID NO: 35)	TTAAGCGGGCGTTGAT (SEQ ID NO: 804)
ONECUT2 (SEQ ID NO: 35)	TTAAGTGGGTGTTGAT (SEQ ID NO: 805)
ONECUT2 (SEQ ID NO: 35)	TAGAGGCGCGGGTTAT (SEQ ID NO: 806)
ONECUT2 (SEQ ID NO: 35)	TAGAGGTGTGGGTTAT (SEQ ID NO: 807)
BCL6 (SEQ ID NO: 36)	ATTTGAAATATGTCGG (SEQ ID NO: 1004)
BCL6	ATTTTGAATATGTTGCT

Gene	Oligo:
(SEQ ID NO: 36)	(SEQ ID NO: 1005)
BCL6 (SEQ ID NO: 36)	ATTCGAGACGTTTTGT (SEQ ID NO: 1006)
BCL6 (SEQ ID NO: 36)	TTTGAGATGTTTTGTTTA (SEQ ID NO: 1007)
BCL6 (SEQ ID NO: 36)	TTCGAGTTTCGAATCGG (SEQ ID NO: 1008)
BCL6 (SEQ ID NO: 36)	TTTGAGTTTTGAATTGGA (SEQ ID NO: 1009)
BCL6 (SEQ ID NO: 36)	ATAGCGAAGGCGTCGA (SEQ ID NO: 1010)
BCL6 (SEQ ID NO: 36)	TATAGTGAAGGTGTTGA (SEQ ID NO: 1011)
WBP11 (SEQ ID NO: 37)	TTACGAGAAGCGGGTA (SEQ ID NO: 946)
WBP11 (SEQ ID NO: 37)	ATTATGAGAAGTGGGTA (SEQ ID NO: 947)
WBP11 (SEQ ID NO: 37)	AGGGGGCGATTTTCGG (SEQ ID NO: 948)
WBP11 (SEQ ID NO: 37)	TAGGGGGTGATTTTGG (SEQ ID NO: 949)
WBP11 (SEQ ID NO: 37)	TTAGCGTCGTTTGATT (SEQ ID NO: 950)
WBP11 (SEQ ID NO: 37)	TTT TAGTGTTGTTTGATT (SEQ ID NO: 951)

Gene	Oligo:
WBP11 (SEQ ID NO: 37)	AGTTCGTTTTATTGCGT (SEQ ID NO: 952)
WBP11 (SEQ ID NO: 37)	GAGTTTGTTTTATTGTGT (SEQ ID NO: 953)
(MX1) (SEQ ID NO: 38)	AACGCGCGAAAGTAAA (SEQ ID NO: 576)
(MX1) (SEQ ID NO: 38)	TTGGGAATGTGTGAAA (SEQ ID NO: 577)
(MX1) (SEQ ID NO: 38)	TTGAGTTGGGTCGAGA (SEQ ID NO: 578)
(MX1) (SEQ ID NO: 38)	TTTGAGTTGGGTTGAGA (SEQ ID NO: 579)
(MX1) (SEQ ID NO: 38)	TATGCGCGGGAAGATT (SEQ ID NO: 580)
(MX1) (SEQ ID NO: 38)	GTATGTGTGGGAAGAT (SEQ ID NO: 581)
(MX1) (SEQ ID NO: 38)	ATTTACGGTTGCGCGG (SEQ ID NO: 582)
(MX1) (SEQ ID NO: 38)	TATGGTTGTGTGGGTTA (SEQ ID NO: 583)
N.N. (SEQ ID NO: 39)	AGGCGTTTATAGTCGGT (SEQ ID NO: 584)
N.N. (SEQ ID NO: 39)	AGGTGTTTATAGTTGGT (SEQ ID NO: 585)
N.N. (SEQ ID NO: 39)	TTTCGAGTTCGGAGTA (SEQ ID NO: 586)

Gene	Oligo:
39)	
N.N. (SEQ ID NO: 39)	TTTTGAGTTTGGAGTAG (SEQ ID NO: 587)
N.N. (SEQ ID NO: 39)	TTGTCGGTCGTAGCGG (SEQ ID NO: 588)
N.N. (SEQ ID NO: 39)	TTTGTTGGTTGTAGTGG (SEQ ID NO: 589)
N.N. (SEQ ID NO: 39)	TTCGTTACGGCGGTAG (SEQ ID NO: 590)
N.N. (SEQ ID NO: 39)	AGTTTGTTATGGTGGT (SEQ ID NO: 591)
APP (SEQ ID NO: 40)	TGAAACGAGGCGGAGA (SEQ ID NO: 592)
APP (SEQ ID NO: 40)	TGAAATGAGGTGGAGA (SEQ ID NO: 593)
APP (SEQ ID NO: 40)	GACGTTGCGTTTTTCGG (SEQ ID NO: 594)
APP (SEQ ID NO: 40)	GGATGTTGTGTTTTTGG (SEQ ID NO: 595)
APP (SEQ ID NO: 40)	TTTTTAGCGGGTCGGA (SEQ ID NO: 596)
APP (SEQ ID NO: 40)	TTTTTAGTGGGTGGA (SEQ ID NO: 597)
APP (SEQ ID NO: 40)	GGACGTTGTAAGCGG (SEQ ID NO: 598)
APP	GGATGTTTGTAAGTGG

Gene	Oligo:
(SEQ ID NO: 40)	(SEQ ID NO: 599)
ORC4L (SEQ ID NO: 41)	TTATACGCGTTGTTTAT (SEQ ID NO: 600)
ORC4L (SEQ ID NO: 41)	TGTATTATATGTGTTGTTT (SEQ ID NO: 601)
ORC4L (SEQ ID NO: 41)	AGCGTGACGGTTCGAG (SEQ ID NO: 602)
ORC4L (SEQ ID NO: 41)	AGTGTGATGGTTTGAG (SEQ ID NO: 603)
ORC4L (SEQ ID NO: 41)	ATTAGGCGAGTTTCGT (SEQ ID NO: 604)
ORC4L (SEQ ID NO: 41)	TTAGGTGAGTTTTGTTT (SEQ ID NO: 605)
NETO1 (SEQ ID NO: 42)	TACGTTTCGGTTTTACGA (SEQ ID NO: 606)
NETO1 (SEQ ID NO: 42)	TTATGTTTGGTTTTATGAT (SEQ ID NO: 607)
NETO1 (SEQ ID NO: 42)	TTACGTCGGTTTCGAT (SEQ ID NO: 608)
NETO1 (SEQ ID NO: 42)	TTTATGTTGGTTTTGATT (SEQ ID NO: 609)
NETO1 (SEQ ID NO: 42)	TTCGGTTTCGGGAAAG (SEQ ID NO: 610)
NETO1 (SEQ ID NO: 42)	TTTGGTTTTGGGAAAGG (SEQ ID NO: 611)

Gene	Oligo:
NETO1 (SEQ ID NO: 42)	TGTCGTACGTGTTTAT (SEQ ID NO: 612)
NETO1 (SEQ ID NO: 42)	AATTTTGTGTATGTGT (SEQ ID NO: 613)
TBC1D3 (SEQ ID NO: 43)	TATTCGCGGGCGGTTT (SEQ ID NO: 988)
TBC1D3 (SEQ ID NO: 43)	TAGTATTTGTGGGTGG (SEQ ID NO: 989)
TBC1D3 (SEQ ID NO: 43)	ATTCGGCGGGAGATTA (SEQ ID NO: 990)
TBC1D3 (SEQ ID NO: 43)	AGTAAATTTGGTGGGA (SEQ ID NO: 991)
TBC1D3 (SEQ ID NO: 43)	AGATTAGTCGAAAGAGT (SEQ ID NO: 992)
TBC1D3 (SEQ ID NO: 43)	GAGATTAGTTGAAAGAGT (SEQ ID NO: 993)
TBC1D3 (SEQ ID NO: 43)	TATATTTTCGGGGTTTAA (SEQ ID NO: 994)
TBC1D3 (SEQ ID NO: 43)	TATATTTTGGGGTTTAAA (SEQ ID NO: 995)
GRB7 (SEQ ID NO: 44)	ATAGTTTCGTTATTTGTAT (SEQ ID NO: 1062)
GRB7 (SEQ ID NO: 44)	GGTATAGTTTTGTTATTTG (SEQ ID NO: 1063)
GRB7 (SEQ ID NO: 44)	TTTAGTACGGGGTGTA (SEQ ID NO: 1064)

Gene	Oligo:
44)	
GRB7 (SEQ ID NO: 44)	TTTTAGTATGGGGTGTA (SEQ ID NO: 1065)
GRB7 (SEQ ID NO: 44)	GGCGTTATAGTTACGTTT (SEQ ID NO: 1066)
GRB7 (SEQ ID NO: 44)	GGGTGTTATAGTTATGTT (SEQ ID NO: 1067)
GRB7 (SEQ ID NO: 44)	TGTTTATCGAAGGTAGA (SEQ ID NO: 1068)
GRB7 (SEQ ID NO: 44)	TGTTTATTGAAGGTAGAA (SEQ ID NO: 1069)
CYP2D6 (SEQ ID NO: 45)	GAGATCGCGTTTTCGT (SEQ ID NO: 844)
CYP2D6 (SEQ ID NO: 45)	AGAGATTGTGTTTTTGT (SEQ ID NO: 845)
CYP2D6 (SEQ ID NO: 45)	ATTCGCGGCGAGGATA (SEQ ID NO: 846)
CYP2D6 (SEQ ID NO: 45)	GATTTGTGGTGAGGAT (SEQ ID NO: 847)
CYP2D6 (SEQ ID NO: 45)	GTCGTTTCGGGGACGT (SEQ ID NO: 848)
CYP2D6 (SEQ ID NO: 45)	GTTGTTTTGGGGATGTG (SEQ ID NO: 849)
CYP2D6 (SEQ ID NO: 45)	TAAGTAGCGTCGATAG (SEQ ID NO: 850)
CYP2D6	AAGTAGTGTTGATAGGG

Gene	Oligo:
(SEQ ID NO: 45)	(SEQ ID NO: 851)
CDK6 (SEQ ID NO: 46)	TACGAATGCGTGGCGG (SEQ ID NO: 866)
CDK6 (SEQ ID NO: 46)	TATGAATGTGTGGTGA (SEQ ID NO: 867)
CDK6 (SEQ ID NO: 46)	TTTCGGAGTAGGCGAG (SEQ ID NO: 868)
CDK6 (SEQ ID NO: 46)	TTTTGGAGTAGGTGAG (SEQ ID NO: 869)
CDK6 (SEQ ID NO: 46)	TACGTTAGTTTCGCGG (SEQ ID NO: 870)
CDK6 (SEQ ID NO: 46)	TATGTTAGTTTTGTGGG (SEQ ID NO: 871)
CDK6 (SEQ ID NO: 46)	ATTGAGACGCGTTTGG (SEQ ID NO: 872)
CDK6 (SEQ ID NO: 46)	GAGATGTGTTTGGGTA (SEQ ID NO: 873)
(Chr. 1p13.2) (SEQ ID NO: 47)	TAAATTCGACGGGTTT (SEQ ID NO: 1054)
(Chr. 1p13.2) (SEQ ID NO: 47)	ATTTGATGGGTTTTTGT (SEQ ID NO: 1055)
(Chr. 1p13.2) (SEQ ID NO: 47)	TTTTCGTTCGGCGGAG (SEQ ID NO: 1056)
(Chr. 1p13.2) (SEQ ID NO: 47)	TTTGTTTGGTGGAGGTT (SEQ ID NO: 1057)

Gene	Oligo:
(Chr. 1p13.2) (SEQ ID NO: 47)	TTCGCGTTTATCGTGT (SEQ ID NO: 1058)
(Chr. 1p13.2) (SEQ ID NO: 47)	TGGTTTGTGTTTATTGT (SEQ ID NO: 1059)
(Chr. 1p13.2) (SEQ ID NO: 47)	TTTCGCGGTTCGTAGT (SEQ ID NO: 1060)
(Chr. 1p13.2) (SEQ ID NO: 47)	TTTGTGGTTTGTAGTTTA (SEQ ID NO: 1061)
(Chr. 17q25.1) (SEQ ID NO: 48)	TTAGGTCGGGAGGAAA (SEQ ID NO: 614)
(Chr. 17q25.1) (SEQ ID NO: 48)	TTAGGTTGGGAGGAAA (SEQ ID NO: 615)
(Chr. 17q25.1) (SEQ ID NO: 48)	TTAGACGTGGGGCGAT (SEQ ID NO: 616)
(Chr. 17q25.1) (SEQ ID NO: 48)	TTAGATGTGGGGTGAT (SEQ ID NO: 617)
(Chr. 17q25.1) (SEQ ID NO: 48)	TAAGGTACGAGCGTGT (SEQ ID NO: 618)
(Chr. 17q25.1) (SEQ ID NO: 48)	AAGGTATGAGTGTGTG (SEQ ID NO: 619)
(Chr. 17q25.1) (SEQ ID NO: 48)	GTAGAGTACGAGAGATT (SEQ ID NO: 620)
(Chr. 17q25.1) (SEQ ID NO: 48)	GGTAGAGTATGAGAGAT (SEQ ID NO: 621)
ABCA8 (SEQ ID NO:)	ATTTGGTTTCGAAGTTT (SEQ ID NO: 996)

Gene	Oligo:
49)	
ABCA8 (SEQ ID NO: 49)	TATTTGGTTTTGAAGTTT (SEQ ID NO: 997)
ABCA8 (SEQ ID NO: 49)	TTTTCGGAATTCGGGT (SEQ ID NO: 998)
ABCA8 (SEQ ID NO: 49)	TTTTGGAATTTGGGTGT (SEQ ID NO: 999)
ABCA8 (SEQ ID NO: 49)	TTTCGGTTTTTAACGGT (SEQ ID NO: 1000)
ABCA8 (SEQ ID NO: 49)	TTTTGGTTTTTAATGGTG (SEQ ID NO: 1001)
ABCA8 (SEQ ID NO: 49)	AAAATTTACGAGGGGA (SEQ ID NO: 1002)
ABCA8 (SEQ ID NO: 49)	TTAAAATTTATGAGGGGA (SEQ ID NO: 1003)
(Chr. 12q14.3) (SEQ ID NO: 50)	ATGACGATGATTGGCGA (SEQ ID NO: 622)
(Chr. 12q14.3) (SEQ ID NO: 50)	GATGATGATTGGTGAGT (SEQ ID NO: 623)
(Chr. 12q14.3) (SEQ ID NO: 50)	TTATGACGTTTAATCGT (SEQ ID NO: 624)
(Chr. 12q14.3) (SEQ ID NO: 50)	AGTTATGATGTTTAATTGT (SEQ ID NO: 625)
(Chr. 12q14.3) (SEQ ID NO: 50)	AATCGAACGTTGGCGT (SEQ ID NO: 626)
(Chr. 12q14.3)	AAATTGAATGTTGGTGT

Gene	Oligo:
(SEQ ID NO: 50)	(SEQ ID NO: 627)
(Chr. 8q12.1) (SEQ ID NO: 51)	TATTCGGGTTTCGCGA (SEQ ID NO: 1070)
(Chr. 8q12.1) (SEQ ID NO: 51)	ATTTGGGTTTGTGAG (SEQ ID NO: 1071)
(Chr. 8q12.1) (SEQ ID NO: 51)	TATTGTTACGCGTCGA (SEQ ID NO: 1072)
(Chr. 8q12.1) (SEQ ID NO: 51)	ATTGTTATGTGTTGATTT (SEQ ID NO: 1073)
(Chr. 8q12.1) (SEQ ID NO: 51)	GACGTGTAGGTCGTAT (SEQ ID NO: 1074)
(Chr. 8q12.1) (SEQ ID NO: 51)	GATGTGTAGGTTGTATT (SEQ ID NO: 1075)
(Chr. 8q12.1) (SEQ ID NO: 51)	TTCGGGAACGATTTTT (SEQ ID NO: 1076)
(Chr. 8q12.1) (SEQ ID NO: 51)	GGGTTTGGGAATGATT (SEQ ID NO: 1077)
MARK2 (SEQ ID NO: 52)	ATATTTTCGGGGGAAGT (SEQ ID NO: 628)
MARK2 (SEQ ID NO: 52)	TATATTTTGGGGGAAGT (SEQ ID NO: 629)
MARK2 (SEQ ID NO: 52)	TTTCGTATTTGTCGGA (SEQ ID NO: 630)
MARK2 (SEQ ID NO: 52)	TTTGTATTTGTTGGAGT (SEQ ID NO: 631)

Gene	Oligo:
MARK2 (SEQ ID NO: 52)	GGTTATATCGTAGGGTA (SEQ ID NO: 632)
MARK2 (SEQ ID NO: 52)	GGGTTATATTGTAGGGT (SEQ ID NO: 633)
MARK2 (SEQ ID NO: 52)	AGGGGGACGAATTAGG (SEQ ID NO: 634)
MARK2 (SEQ ID NO: 52)	GAGGGGGATGAATTAG (SEQ ID NO: 635)
ELK1 (SEQ ID NO: 53)	GGTCGGCGTTGATTTTA (SEQ ID NO: 920)
ELK1 (SEQ ID NO: 53)	GGTTGGTGTTGATTTTA (SEQ ID NO: 921)
ELK1 (SEQ ID NO: 53)	GTCGGGATTCGAACGG (SEQ ID NO: 922)
ELK1 (SEQ ID NO: 53)	GTTGGGATTTGAATGG (SEQ ID NO: 923)
ELK1 (SEQ ID NO: 53)	GTCGGAAGTTTCGGGA (SEQ ID NO: 924)
ELK1 (SEQ ID NO: 53)	GTTGGAAGTTTGTGGAT (SEQ ID NO: 925)
ELK1 (SEQ ID NO: 53)	ATATCGTAGGGTAGGCGG (SEQ ID NO: 926)
ELK1 (SEQ ID NO: 53)	ATATTGTAGGGTAGGTGG (SEQ ID NO: 927)
Q8WUT3 (SEQ ID NO:	TAGAACGGCGTGGGAT (SEQ ID NO: 636)

Gene	Oligo:
54)	
Q8WUT3 (SEQ ID NO: 54)	TAGAATGGTGTGGGAT (SEQ ID NO: 637)
Q8WUT3 (SEQ ID NO: 54)	GTCGCGATGTAGTTACGT (SEQ ID NO: 638)
Q8WUT3 (SEQ ID NO: 54)	GTTGTGATGTAGTTATGT (SEQ ID NO: 639)
Q8WUT3 (SEQ ID NO: 54)	TTAGTTTCGGGATCGG (SEQ ID NO: 640)
Q8WUT3 (SEQ ID NO: 54)	TTTAGTTTTTGGGATTGG (SEQ ID NO: 641)
Q8WUT3 (SEQ ID NO: 54)	TTCGTTTTTCGGGATA (SEQ ID NO: 642)
Q8WUT3 (SEQ ID NO: 54)	TTTGTTTTTTGGGATAAA (SEQ ID NO: 643)
CGB (SEQ ID NO: 55)	TTACGTCGTGGTTTTTA (SEQ ID NO: 954)
CGB (SEQ ID NO: 55)	TTATGTTGTGGTTTTTAG (SEQ ID NO: 955)
CGB (SEQ ID NO: 55)	GGCGTGAATTCGTGG (SEQ ID NO: 956)
CGB (SEQ ID NO: 55)	GGTGTGAATTTTGTGGT (SEQ ID NO: 957)
CGB (SEQ ID NO: 55)	TTTCGAGTTTATTCGGT (SEQ ID NO: 958)
CGB	TTTTGAGTTTATTTGGTT

Gene	Oligo:
(SEQ ID NO: 55)	(SEQ ID NO: 959)
CGB (SEQ ID NO: 55)	TTATCGCGATGTGCGT (SEQ ID NO: 960)
CGB (SEQ ID NO: 55)	ATTATTGTGATGTGTGT (SEQ ID NO: 961)
BSG (SEQ ID NO: 56)	TACGGTTCGCGTTGTT (SEQ ID NO: 644)
BSG (SEQ ID NO: 56)	GGAGTATGGTTTGTGT (SEQ ID NO: 645)
BSG (SEQ ID NO: 56)	GTAAGGTTTCGGCGAGA (SEQ ID NO: 646)
BSG (SEQ ID NO: 56)	GTAAGGTTTGGTGAGA (SEQ ID NO: 647)
BSG (SEQ ID NO: 56)	TTACGTTTTTCGGGAAG (SEQ ID NO: 648)
BSG (SEQ ID NO: 56)	TTATGTTTTTGGGAAGG (SEQ ID NO: 649)
BSG (SEQ ID NO: 56)	TACGTTTCGAGGATCGG (SEQ ID NO: 650)
BSG (SEQ ID NO: 56)	TATGTTTTGAGGATTGG (SEQ ID NO: 651)
BCKDK (SEQ ID NO: 57)	GGGCGTTAGGCGGATT (SEQ ID NO: 652)
BCKDK (SEQ ID NO: 57)	TGGGTGTTAGGTGGAT (SEQ ID NO: 653)

Gene	Oligo:
BCKDK (SEQ ID NO: 57)	AGAGCGGTTAGCGTAG (SEQ ID NO: 654)
BCKDK (SEQ ID NO: 57)	TGAGAGTGGTTAGTGT (SEQ ID NO: 655)
BCKDK (SEQ ID NO: 57)	ATAGAGGGCGTGAATT (SEQ ID NO: 656)
BCKDK (SEQ ID NO: 57)	AGAGGGTGTGAATTTT (SEQ ID NO: 657)
BCKDK (SEQ ID NO: 57)	TAGGATTTACGAGGAAA (SEQ ID NO: 658)
BCKDK (SEQ ID NO: 57)	AGGATTTATGAGGAAAAT (SEQ ID NO: 659)
SOX8 (SEQ ID NO: 58)	TTTTCGGTTCGAAGTA (SEQ ID NO: 660)
SOX8 (SEQ ID NO: 58)	TTTTGGTTTGAAGTAGG (SEQ ID NO: 661)
SOX8 (SEQ ID NO: 58)	AGGTCGTTTTTATCGA (SEQ ID NO: 662)
SOX8 (SEQ ID NO: 58)	AGGTTGTTTTTATTGAGT (SEQ ID NO: 663)
SOX8 (SEQ ID NO: 58)	GTAGTTACGGGGCGTT (SEQ ID NO: 664)
SOX8 (SEQ ID NO: 58)	GTAGTTATGGGGTGTT (SEQ ID NO: 665)
SOX8 (SEQ ID NO: 58)	TGTCGTATAGGCGGTT (SEQ ID NO: 666)

Gene	Oligo:
58)	
SOX8 (SEQ ID NO: 58)	TTGTTGTATAGGTGGTT (SEQ ID NO: 667)
DAG1 (SEQ ID NO: 59)	TTTCGTGGCGGAGAAT (SEQ ID NO: 820)
DAG1 (SEQ ID NO: 59)	TTTTGTGGTGGAGAAT (SEQ ID NO: 821)
DAG1 (SEQ ID NO: 59)	TACGGATATTTTCGGTT (SEQ ID NO: 822)
DAG1 (SEQ ID NO: 59)	AATTATGGATATTTTGGTT (SEQ ID NO: 823)
DAG1 (SEQ ID NO: 59)	TTACGATTCGTAGGTT (SEQ ID NO: 824)
DAG1 (SEQ ID NO: 59)	TATTATTATGATTTGTAGGT (SEQ ID NO: 825)
SEMA4B (SEQ ID NO: 60)	AGTTTTGGGCGCGATTT (SEQ ID NO: 668)
SEMA4B (SEQ ID NO: 60)	AGTTTTGGGTGTGATTT (SEQ ID NO: 669)
SEMA4B (SEQ ID NO: 60)	AGCGAATAGATTGCGGAT (SEQ ID NO: 670)
SEMA4B (SEQ ID NO: 60)	AGTGAATAGATTGTGGAT (SEQ ID NO: 671)
SEMA4B (SEQ ID NO: 60)	AGCGATTAGATTGCGGAT (SEQ ID NO: 672)
SEMA4B	AGTGATTAGATTGTGGAT

Gene	Oligo:
(SEQ ID NO: 60)	(SEQ ID NO: 673)
SEMA4B (SEQ ID NO: 60)	TAGGCGTTCGATTTTT (SEQ ID NO: 674)
SEMA4B (SEQ ID NO: 60)	GGGTAGGTGTTTGATT (SEQ ID NO: 675)
APC (SEQ ID NO: 2)	GGTTTCGTTTAATCGT (SEQ ID NO: 928)
APC (SEQ ID NO: 2)	GGGTTTTGTTTAATTGTA (SEQ ID NO: 929)
APC (SEQ ID NO: 2)	PTCGTATTAGCGGAT (SEQ ID NO: 930)
APC (SEQ ID NO: 2)	GGTTTGTATTTAGTGGA (SEQ ID NO: 931)
APC (SEQ ID NO: 2)	ATCGGCGGGTTTTCGA (SEQ ID NO: 932)
APC (SEQ ID NO: 2)	AATTGGTGGGTTTTTGA (SEQ ID NO: 933)
APC (SEQ ID NO: 2)	ATTTTCGAGTTCGGTA (SEQ ID NO: 934)
APC (SEQ ID NO: 2)	TTTTTGAGTTTGGTAGT (SEQ ID NO: 935)
CDKN2A (SEQ ID NO: 3)	GGCGTTGTTTAACGTAT (SEQ ID NO: 676)
CDKN2A (SEQ ID NO: 3)	GGGTGTTGTTTAATGTA (SEQ ID NO: 677)
CDKN2A (SEQ ID NO: 3)	AACGTATCGAATAGTTACGG (SEQ ID NO: 678)
CDKN2A (SEQ ID NO: 3)	AATGTATTGAATAGTTATGG (SEQ ID NO: 679)
CDKN2A (SEQ ID NO: 3)	TACGGTCGGAGGTCGA (SEQ ID NO: 680)
CDKN2A (SEQ ID NO: 3)	TATGGTTGGAGGTTGA (SEQ ID NO: 681)
CSPG2 (SEQ ID NO: 4)	TTCGGTTAGTTTCGTAT (SEQ ID NO: 904)

Gene	Oligo:
CSPG2 (SEQ ID NO: 4)	TTTTGGTTAGTTTTGTATT (SEQ ID NO: 905)
CSPG2 (SEQ ID NO: 4)	TTCGGGTTATTACGTTT (SEQ ID NO: 906)
CSPG2 (SEQ ID NO: 4)	TTTTGGGTTATTATGTTTT (SEQ ID NO: 907)
CSPG2 (SEQ ID NO: 4)	TTTAGTCGCGTAGCGT (SEQ ID NO: 908)
CSPG2 (SEQ ID NO: 4)	ATTTAGTTGTGTAGTGTT (SEQ ID NO: 909)
CSPG2 (SEQ ID NO: 4)	AATTCGCGAGTTTAGA (SEQ ID NO: 910)
CSPG2 (SEQ ID NO: 4)	GAAAAAAATTTGTGAGTT (SEQ ID NO: 911)
ERBB2 (SEQ ID NO: 5)	TGTGAGAACGTTGTA (SEQ ID NO: 912)
ERBB2 (SEQ ID NO: 5)	TGAGAATGGTTGTAGG (SEQ ID NO: 913)
ERBB2 (SEQ ID NO: 5)	TTAGGCGTTTCGGCGT (SEQ ID NO: 914)
ERBB2 (SEQ ID NO: 5)	TTTAGGTGTTTTGGTGT (SEQ ID NO: 915)
ERBB2 (SEQ ID NO: 5)	TAGGTTTGCGCGAAGA (SEQ ID NO: 916)
ERBB2 (SEQ ID NO: 5)	TTTGTGTGAAGAGAGG (SEQ ID NO: 917)
ERBB2 (SEQ ID NO: 5)	TAATTATCGGAGAAGGA (SEQ ID NO: 918)
ERBB2 (SEQ ID NO: 5)	TAATTATTGGAGAAGGAG (SEQ ID NO: 919)
STMN1 (SEQ ID NO: 6)	TTAGGCGGTTTCGGATT (SEQ ID NO: 1012)
STMN1 (SEQ ID NO: 6)	TTAGGTGGTTTGGATT (SEQ ID NO: 1013)
STMN1 (SEQ ID NO: 6)	TATCGGTTTCGGGAATT (SEQ ID NO: 1014)
STMN1 (SEQ ID NO: 6)	TATTGGTTTGGGAATTT (SEQ ID NO: 1015)

Gene	Oligo:
STMN1 (SEQ ID NO: 6)	TTTCGCGCGGAGGTTA (SEQ ID NO: 1016)
STMN1 (SEQ ID NO: 6)	TTTTGTGTGGAGGTTA (SEQ ID NO: 1017)
STMN1 (SEQ ID NO: 6)	GGTAAGAACGTATATAGT (SEQ ID NO: 1018)
STMN1 (SEQ ID NO: 6)	TGGTAAGAATGTATATAGT (SEQ ID NO: 1019)
STMN1 (SEQ ID NO: 6)	TTTCGGTTAATGCGGA (SEQ ID NO: 1020)
STMN1 (SEQ ID NO: 6)	TTTTTGGTTAATGTGGA (SEQ ID NO: 1021)
STMN1 (SEQ ID NO: 6)	TACGTTGCGGATTTGT (SEQ ID NO: 1022)
STMN1 (SEQ ID NO: 6)	AGGGTTATGTTTGTGA (SEQ ID NO: 1023)
STMN1 (SEQ ID NO: 6)	GATACGTCGGTGTCGG (SEQ ID NO: 1024)
STMN1 (SEQ ID NO: 6)	TGATATGTTGGTGTGG (SEQ ID NO: 1025)
STMN1 (SEQ ID NO: 6)	TTACGGCGAGATTATT (SEQ ID NO: 1026)
STMN1 (SEQ ID NO: 6)	TTTTATGGTGAGATTATT (SEQ ID NO: 1027)
STK11 (SEQ ID NO: 7)	ATTAATCGTCGTTTCGG (SEQ ID NO: 880)
STK11 (SEQ ID NO: 7)	GATTAATTGTTGTTTGGG (SEQ ID NO: 881)
STK11 (SEQ ID NO: 7)	TAATCGTTAGCGGCGG (SEQ ID NO: 882)
STK11 (SEQ ID NO: 7)	TTAATTGTTAGTGGTGG (SEQ ID NO: 883)
STK11 (SEQ ID NO: 7)	GTCGTTTTTCGCGAGGA (SEQ ID NO: 884)
STK11 (SEQ ID NO: 7)	GTTGTTTTTGTGAGGAG (SEQ ID NO: 885)
STK11 (SEQ ID NO: 7)	TAATGAGCGCGTTGTA (SEQ ID NO: 886)

Gene	Oligo:
STK11 (SEQ ID NO: 7)	ATGAGTGTGTTGTATTT (SEQ ID NO: 887)
CA9 (SEQ ID NO: 8)	ATGGTTTCGATAATTTTT (SEQ ID NO: 682)
CA9 (SEQ ID NO: 8)	ATGGTTTTGATAATTTTTT (SEQ ID NO: 683)
CA9 (SEQ ID NO: 8)	TGTACGTATAGTTCGTA (SEQ ID NO: 684)
CA9 (SEQ ID NO: 8)	TTAATGTATGTATAGTTTGT (SEQ ID NO: 685)
CA9 (SEQ ID NO: 8)	ATATATCGTGTGTTGGG (SEQ ID NO: 686)
CA9 (SEQ ID NO: 8)	ATATATTGTGTGTTGGG (SEQ ID NO: 687)
CA9 (SEQ ID NO: 8)	ATAGTTAGTCGTATGGT (SEQ ID NO: 688)
CA9 (SEQ ID NO: 8)	ATAGTTAGTTGTATGGTT (SEQ ID NO: 689)
PAX6 (SEQ ID NO: 9)	TATTGTTTCGGTTGTTAG (SEQ ID NO: 690)
PAX6 (SEQ ID NO: 9)	TATTGTTTTGGTTGTTAG (SEQ ID NO: 691)
PAX6 (SEQ ID NO: 9)	GGCGACGCGGTTAGTT (SEQ ID NO: 692)
PAX6 (SEQ ID NO: 9)	GGTGATGTGGTTAGTT (SEQ ID NO: 693)
PAX6 (SEQ ID NO: 9)	TAGGTCGCGTAGATTT (SEQ ID NO: 694)
PAX6 (SEQ ID NO: 9)	AGTTTAGGTTGTGTAGA (SEQ ID NO: 695)
PAX6 (SEQ ID NO: 9)	TAGCGTATTTTTCGGT (SEQ ID NO: 696)
PAX6 (SEQ ID NO: 9)	TAGTGTATTTTTTGGTTG (SEQ ID NO: 697)
SFN (SEQ ID NO: 10)	AGTAGGTCGAACGTTA (SEQ ID NO: 698)
SFN	AGAGTAGGTTGAATGTT

Gene	Oligo:
(SEQ ID NO: 10)	(SEQ ID NO: 699)
SFN (SEQ ID NO: 10)	TTGCGAAGAGCGAAAT (SEQ ID NO: 700)
SFN (SEQ ID NO: 10)	TGTGAAGAGTGAAATTT (SEQ ID NO: 701)
SFN (SEQ ID NO: 10)	TTTCGAGGTGCGTGAGT (SEQ ID NO: 702)
SFN (SEQ ID NO: 10)	TTTGAGGTGTGTGAGTA (SEQ ID NO: 703)
SFN (SEQ ID NO: 10)	TGTGCGATATCGTGTT (SEQ ID NO: 704)
SFN (SEQ ID NO: 10)	TGTGATATTGTGTTGGG (SEQ ID NO: 705)
S100A2 (SEQ ID NO: 11)	TTTAATTGCGGTTGTGTG (SEQ ID NO: 786)
S100A2 (SEQ ID NO: 11)	TTTAATTGTGGTTGTGTG (SEQ ID NO: 787)
S100A2 (SEQ ID NO: 11)	TATATAGGCGTATGTATG (SEQ ID NO: 788)
S100A2 (SEQ ID NO: 11)	TATATAGGTGTATGTATG (SEQ ID NO: 789)
S100A2 (SEQ ID NO: 11)	TGTATACGAGTATTGGA (SEQ ID NO: 790)
S100A2 (SEQ ID NO: 11)	TATGTATATGAGTATTGGA (SEQ ID NO: 791)

Gene	Oligo:
S100A2 (SEQ ID NO: 11)	AGTTTTAGCGTGTGTTTA (SEQ ID NO: 792)
S100A2 (SEQ ID NO: 11)	AGTTTTAGTGTGTGTTTA (SEQ ID NO: 793)
TFF1 (SEQ ID NO: 12)	AGAATTTATCGTATAAAAAG (SEQ ID NO: 794)
TFF1 (SEQ ID NO: 12)	AATTTATTGTATAAAAAGGT (SEQ ID NO: 795)
TFF1 (SEQ ID NO: 12)	GGACGTCGATGGTATT (SEQ ID NO: 796)
TFF1 (SEQ ID NO: 12)	AGGGATGTTGATGGTA (SEQ ID NO: 797)
TFF1 (SEQ ID NO: 12)	AACGGTGTCGTCGAAA (SEQ ID NO: 798)
TFF1 (SEQ ID NO: 12)	AATGGTGTTGTTGAAAT (SEQ ID NO: 799)
TGFBR2 (SEQ ID NO: 13)	AAAACGTGGACGTTTT (SEQ ID NO: 896)
TGFBR2 (SEQ ID NO: 13)	GAAAATGTGGATGTTTT (SEQ ID NO: 897)
TGFBR2 (SEQ ID NO: 13)	TGAAAGTCGGTTAAAGT (SEQ ID NO: 898)
TGFBR2 (SEQ ID NO: 13)	TGAAAGTTGGTTAAAGT (SEQ ID NO: 899)
TGFBR2 (SEQ ID NO:	TTGGACGTCGAGGAGA (SEQ ID NO: 900)

Gene	Oligo:
13)	
TGFBR2 (SEQ ID NO: 13)	TTGGATGTTGAGGAGA (SEQ ID NO: 901)
TGFBR2 (SEQ ID NO: 13)	TTTTCGGGCGGAGAGA (SEQ ID NO: 902)
TGFBR2 (SEQ ID NO: 13)	AAGGTTTTTGGGTGGA (SEQ ID NO: 903)
TP53 (SEQ ID NO: 14)	TATTAGGTCGGCGAGA (SEQ ID NO: 858)
TP53 (SEQ ID NO: 14)	AGGTTGGTGAGAATTT (SEQ ID NO: 859)
TP53 (SEQ ID NO: 14)	TTCGGTAGGCGGATTA (SEQ ID NO: 860)
TP53 (SEQ ID NO: 14)	TTTTTGGTAGGTGGAT (SEQ ID NO: 861)
TP53 (SEQ ID NO: 14)	ATATTTTGC GTTCGGG (SEQ ID NO: 862)
TP53 (SEQ ID NO: 14)	ATATTTTGTGTTTGGGT (SEQ ID NO: 863)
TP53 (SEQ ID NO: 14)	TACGACGGTGATACGT (SEQ ID NO: 864)
TP53 (SEQ ID NO: 14)	TTTATGATGGTGATATGT (SEQ ID NO: 865)
TP73 (SEQ ID NO: 15)	TTCGTTCGCGAAGTTA (SEQ ID NO: 706)
TP73	GGTTTGTTTGTGAAGTTA

Gene	Oligo:
(SEQ ID NO: 15)	(SEQ ID NO: 707)
PLAU (SEQ ID NO: 16)	AAGAGGTCGTCGGGAT (SEQ ID NO: 708)
PLAU (SEQ ID NO: 16)	AAGAGGTTGTTGGGAT (SEQ ID NO: 709)
PLAU (SEQ ID NO: 16)	TTATCGCGGGTATTTT (SEQ ID NO: 710)
PLAU (SEQ ID NO: 16)	TTGGTTATTGTGGGTAT (SEQ ID NO: 711)
PLAU (SEQ ID NO: 16)	TTCGATTTCGTTATTATG (SEQ ID NO: 712)
PLAU (SEQ ID NO: 16)	TTTGATTTTGTTATTATGAG (SEQ ID NO: 713)
PLAU (SEQ ID NO: 16)	GTCGTGAGCGATTTTA (SEQ ID NO: 714)
PLAU (SEQ ID NO: 16)	TTGGTTGTGAGTGATT (SEQ ID NO: 715)
TMEFF2 (SEQ ID NO: 17)	TATCGTAGTTCGTTTCGG (SEQ ID NO: 874)
TMEFF2 (SEQ ID NO: 17)	ATTGTAGTTTGTTTGGT (SEQ ID NO: 875)
TMEFF2 (SEQ ID NO: 17)	AAACGTTTATCGGTTG (SEQ ID NO: 876)
TMEFF2 (SEQ ID NO: 17)	AATGTTTATTGGTTGGA (SEQ ID NO: 877)

Gene	Oligo:
TMEFF2 (SEQ ID NO: 17)	TTCGTAGAAGAATACGCGTA (SEQ ID NO: 878)
TMEFF2 (SEQ ID NO: 17)	TTTGTAGAAGAATATGTGTA (SEQ ID NO: 879)
ESR1 (SEQ ID NO: 18)	TGCGGTTGTATACGTAG (SEQ ID NO: 962)
ESR1 (SEQ ID NO: 18)	TGTGTGGTTGTATATGT (SEQ ID NO: 963)
ESR1 (SEQ ID NO: 18)	TTCGTGTTAGATTTTCGATAT (SEQ ID NO: 964)
ESR1 (SEQ ID NO: 18)	TTTGTGTTAGATTTTGATAT (SEQ ID NO: 965)
ESR1 (SEQ ID NO: 18)	AACGCGAAAGACGGAT (SEQ ID NO: 966)
ESR1 (SEQ ID NO: 18)	ATAAATGTGAAAGATGGA (SEQ ID NO: 967)
ESR1 (SEQ ID NO: 18)	GGGCGTACGAGGATTT (SEQ ID NO: 968)
ESR1 (SEQ ID NO: 18)	GGGTGTATGAGGATTT (SEQ ID NO: 969)
HSPB1 (SEQ ID NO: 20)	AGGGTATTCGTCGGTT (SEQ ID NO: 888)
HSPB1 (SEQ ID NO: 20)	AGGGTATTTGTTGGTT (SEQ ID NO: 889)
HSPB1 (SEQ ID NO: 20)	GAATTCGAGAGCGCGA (SEQ ID NO: 892)

Gene	Oligo:
20)	
HSPB1 (SEQ ID NO: 20)	TGAATTTGAGAGTGTGA (SEQ ID NO: 893)
RASSF1 (SEQ ID NO: 21)	AGTAAATCGGATTAGGA (SEQ ID NO: 852)
RASSF1 (SEQ ID NO: 21)	AGTAAATTGGATTAGGAG (SEQ ID NO: 853)
RASSF1 (SEQ ID NO: 21)	TACGGGTATTTTCGCGT (SEQ ID NO: 854)
RASSF1 (SEQ ID NO: 21)	ATATGGGTATTTTGTGT (SEQ ID NO: 855)
RASSF1 (SEQ ID NO: 21)	TGCGAGAGCGCGTTTA (SEQ ID NO: 856)
RASSF1 (SEQ ID NO: 21)	TTGTGAGAGTGTGTTTA (SEQ ID NO: 857)
GRIN2D (SEQ ID NO: 24)	ATTTTCGATTGAGGCGG (SEQ ID NO: 716)
GRIN2D (SEQ ID NO: 24)	ATTTTGATTGAGGTGG (SEQ ID NO: 717)
PSAT1 (SEQ ID NO: 25)	TTCGTCGGTGTTACGT (SEQ ID NO: 718)
PSAT1 (SEQ ID NO: 25)	TTTGTGGTGTTATGT (SEQ ID NO: 719)
PSAT1 (SEQ ID NO: 25)	GGCGAGTTCGGGTAGT (SEQ ID NO: 720)
PSAT1	GGTGAGTTGGGTAGT

Gene	Oligo:
(SEQ ID NO: 25)	(SEQ ID NO: 721)
PSAT1 (SEQ ID NO: 25)	ATAGTAAACGCGAGGA (SEQ ID NO: 818)
PSAT1 (SEQ ID NO: 25)	AGTAAATGTGAGGAGG (SEQ ID NO: 819)
PSAT1 (SEQ ID NO: 25)	AAGTTTTTCGCGAGCGG (SEQ ID NO: 722)
PSAT1 (SEQ ID NO: 25)	AAGTTTTTGTGAGTGG (SEQ ID NO: 723)
PSAT1 (SEQ ID NO: 25)	AGGAAGTTCGGCGAGG (SEQ ID NO: 724)
PSAT1 (SEQ ID NO: 25)	AGGAAGTTTGGTGAGG (SEQ ID NO: 725)
CYP2D6 (SEQ ID NO: 27)	TACGACGATTTTCGTT (SEQ ID NO: 726)
CYP2D6 (SEQ ID NO: 27)	GAGTATGATGATTTTTGT (SEQ ID NO: 727)
CYP2D6 (SEQ ID NO: 27)	TTCGTCGATTAAGTCGG (SEQ ID NO: 728)
CYP2D6 (SEQ ID NO: 27)	TTTGTTGATTAAGTTGGT (SEQ ID NO: 729)
CYP2D6 (SEQ ID NO: 27)	GTGGCGCGAGTAGAGG (SEQ ID NO: 730)
CYP2D6 (SEQ ID NO: 27)	GTGGTGTGAGTAGAGG (SEQ ID NO: 731)

Gene	Oligo:
CYP2D6 (SEQ ID NO: 27)	AACGTTTACGTGTTTCGT (SEQ ID NO: 732)
CYP2D6 (SEQ ID NO: 27)	GTAATGTTTATGTGTTTGT (SEQ ID NO: 733)
COX7A2L (SEQ ID NO: 28)	AATTCGATCGCGGGTA (SEQ ID NO: 1086)
COX7A2L (SEQ ID NO: 28)	ATTTGATTGTGGGTAGA (SEQ ID NO: 1087)
PLAU (SEQ ID NO: 30)	TATTTGTCTCGCTTGAT (SEQ ID NO: 1044)
PLAU (SEQ ID NO: 30)	ATTTGTTGTGTTGATGA (SEQ ID NO: 1045)
PLAU (SEQ ID NO: 30)	TGTAATTCGGGGATTT (SEQ ID NO: 1046)
PLAU (SEQ ID NO: 30)	TTGTAATTTGGGGATTT (SEQ ID NO: 1047)
PLAU (SEQ ID NO: 30)	AGGAAGTACGGAGAAT (SEQ ID NO: 1048)
PLAU (SEQ ID NO: 30)	AGGAAGTATGGAGAATT (SEQ ID NO: 1049)
PLAU (SEQ ID NO: 30)	TTCGTTGGAGATCGCGT (SEQ ID NO: 1050)
PLAU (SEQ ID NO: 30)	TTTGTTGGAGATTGTGT (SEQ ID NO: 1051)
PLAU (SEQ ID NO: 30)	TTGCGGAAGTACGCGG (SEQ ID NO: 1052)

Gene	Oligo:
30)	
PLAU (SEQ ID NO: 30)	TTGTGGAAGTATGTGG (SEQ ID NO: 1053)
VTN (SEQ ID NO: 31)	TTCGGGTTTCGCGAAAG (SEQ ID NO: 1028)
VTN (SEQ ID NO: 31)	TTTGGGTTTGTGAAAG (SEQ ID NO: 1029)
VTN (SEQ ID NO: 31)	TTTTGTTCGCGTTGAA (SEQ ID NO: 1030)
VTN (SEQ ID NO: 31)	TTGTTTGTGTTGAAGTA (SEQ ID NO: 1031)
VTN (SEQ ID NO: 31)	TGGGTCGCGAGGTAGT (SEQ ID NO: 1032)
VTN (SEQ ID NO: 31)	TGGGTTGTGAGGTAGT (SEQ ID NO: 1033)
VTN (SEQ ID NO: 31)	TTCGATGGCGGTTTCGA (SEQ ID NO: 1036)
VTN (SEQ ID NO: 31)	TTTGATGGTGGTTTGA (SEQ ID NO: 1037)
SULT1A1 (SEQ ID NO: 32)	TTCGAGTCGTTTTGAT (SEQ ID NO: 734)
SULT1A1 (SEQ ID NO: 32)	TTTGAGTTGTTTTGATG (SEQ ID NO: 735)
SULT1A1 (SEQ ID NO: 32)	TTCGTCGTGTACGGTT (SEQ ID NO: 736)
SULT1A1	TTTGTGTGTATGGTTT

Gene	Oligo:
(SEQ ID NO: 32)	(SEQ ID NO: 737)
SULT1A1 (SEQ ID NO: 32)	AGGATTTTCGTTTTTCGG (SEQ ID NO: 738)
SULT1A1 (SEQ ID NO: 32)	AGGATTTTGTTTTTTGGG (SEQ ID NO: 739)
SULT1A1 (SEQ ID NO: 32)	TTTTTCGGTTGAAGTCGG (SEQ ID NO: 740)
SULT1A1 (SEQ ID NO: 32)	TTTTTTGGTTGAAGTTGG (SEQ ID NO: 741)
PCAF (SEQ ID NO: 33)	AGCGTCGGTACGTATA (SEQ ID NO: 986)
PCAF (SEQ ID NO: 33)	GGTAGTGTTGGTATGT (SEQ ID NO: 987)
PRKCD (SEQ ID NO: 34)	ATTTTCGCGTTCGGATT (SEQ ID NO: 742)
PRKCD (SEQ ID NO: 34)	GATTTTGTGTTTGGATT (SEQ ID NO: 743)
EGR4 (SEQ ID NO: 1)	AAGCGTATTTATCGGA (SEQ ID NO: 744)
EGR4 (SEQ ID NO: 1)	GGAAGTGTATTTATTGGA (SEQ ID NO: 745)
EGR4 (SEQ ID NO: 1)	TATCGGACGGTCGGTT (SEQ ID NO: 746)
EGR4 (SEQ ID NO: 1)	ATTTATTGGATGGTTGG (SEQ ID NO: 747)
EGR4 (SEQ ID NO: 1)	AGGCGTAGCGTTTTAG (SEQ ID NO: 748)
EGR4 (SEQ ID NO: 1)	TGAGGTGTAGTGTTTT (SEQ ID NO: 749)

Gene	Oligo:
EGR4 (SEQ ID NO: 1)	AACGTTATAGTTCGAGT (SEQ ID NO: 750)
EGR4 (SEQ ID NO: 1)	AATGTTATAGTTTGAGTTT (SEQ ID NO: 751)
TP73 (SEQ ID NO: 15)	GTGCGAGTTAGTCGGA (SEQ ID NO: 752)
TP73 (SEQ ID NO: 15)	GTGTGAGTTAGTTGGA (SEQ ID NO: 753)
TP73 (SEQ ID NO: 15)	TATCGGTTTCGGAGTTA (SEQ ID NO: 754)
TP73 (SEQ ID NO: 15)	AGGATATTGGTTTGGAG (SEQ ID NO: 755)
TP73 (SEQ ID NO: 15)	AGAGTCGTTTCGGAATT (SEQ ID NO: 756)
TP73 (SEQ ID NO: 15)	TGAGAGTTGTTTGGAAAT (SEQ ID NO: 757)
SYK (SEQ ID NO: 19)	GAAGTTATCGCGTTGG (SEQ ID NO: 826)
SYK (SEQ ID NO: 19)	AGAAGTTATTGTGTTGG (SEQ ID NO: 827)
SYK (SEQ ID NO: 19)	GATCGATGCGGTTTAT (SEQ ID NO: 828)
SYK (SEQ ID NO: 19)	GGGATTGATGTGGTTTA (SEQ ID NO: 829)
SYK (SEQ ID NO: 19)	GTTCGGCGGGAGGAGA (SEQ ID NO: 830)
SYK	GTTTGGTGGGAGGAGA

Gene	Oligo:
(SEQ ID NO: 19)	(SEQ ID NO: 831)
SYK (SEQ ID NO: 19)	AGTCGATTTTCGTTTAG (SEQ ID NO: 832)
SYK (SEQ ID NO: 19)	TAGTTGATTTTGTTTAGT (SEQ ID NO: 833)
SYK (SEQ ID NO: 19)	GGAAGAGTCGCGGGTT (SEQ ID NO: 834)
SYK (SEQ ID NO: 19)	GGAAGAGTTGTGGGTT (SEQ ID NO: 835)
HSPB1 (SEQ ID NO: 20)	AGTCGTGTTACGGTAG (SEQ ID NO: 890)
HSPB1 (SEQ ID NO: 20)	AGTTGTGTTATGGTAGG (SEQ ID NO: 891)
HSPB1 (SEQ ID NO: 20)	TTTTTTCGTTAAGGAAAG (SEQ ID NO: 894)
HSPB1 (SEQ ID NO: 20)	TTTTTTTTTGTTAAGGAAAG (SEQ ID NO: 895)
TES (SEQ ID NO: 22)	TAGAAGTCGGTTCGTG (SEQ ID NO: 758)
TES (SEQ ID NO: 22)	AGAAGTTGGTTTGTGG (SEQ ID NO: 759)
TES (SEQ ID NO: 22)	GATTGGGCGGCGGAAG (SEQ ID NO: 760)
TES (SEQ ID NO: 22)	ATTGGGTGGTGGAAAGT (SEQ ID NO: 761)

Gene	Oligo:
TES (SEQ ID NO: 22)	TAGCGGAGTCGGAGGT (SEQ ID NO: 762)
TES (SEQ ID NO: 22)	TAGTGGAGTTGGAGGT (SEQ ID NO: 763)
TES (SEQ ID NO: 22)	AATTCGGTCGTGGGAT (SEQ ID NO: 764)
TES (SEQ ID NO: 22)	AATTTGGTTGTGGGAT (SEQ ID NO: 765)
PITX2 (SEQ ID NO: 23)	AGTCGGGAGAGCGAAA (SEQ ID NO: 970)
PITX2 (SEQ ID NO: 23)	AGTTGGGAGAGTGAAA (SEQ ID NO: 971)
PITX2 (SEQ ID NO: 23)	AAGAGTCGGGAGTCGGA (SEQ ID NO: 972)
PITX2 (SEQ ID NO: 23)	AAGAGTTGGGAGTTGGA (SEQ ID NO: 973)
PITX2 (SEQ ID NO: 23)	GGTCGAAGAGTCGGGA (SEQ ID NO: 974)
PITX2 (SEQ ID NO: 23)	GGTTGAAGAGTTGGGA (SEQ ID NO: 975)
PITX2 (SEQ ID NO: 23)	ATGTTAGCGGGTCGAA (SEQ ID NO: 976)
PITX2 (SEQ ID NO: 23)	TAGTGGGTTGAAGAGT (SEQ ID NO: 977)
GRIN2D (SEQ ID NO: 23)	GAGAGTCGGGATGATT (SEQ ID NO: 766)

Gene	Oligo:
24)	
GRIN2D (SEQ ID NO: 24)	GGAGAGTTGGGATGAT (SEQ ID NO: 767)
GRIN2D (SEQ ID NO: 24)	TAGGGTCGAGATTTGG (SEQ ID NO: 768)
GRIN2D (SEQ ID NO: 24)	TTAGGGTTGAGATTTGG (SEQ ID NO: 769)
GRIN2D (SEQ ID NO: 24)	AGTGTGGCGAATATTG (SEQ ID NO: 770)
GRIN2D (SEQ ID NO: 24)	GTGTGGTGAATATTGAA (SEQ ID NO: 771)
PSAT1 (SEQ ID NO: 25)	TTTCGATTCGGTTTAGA (SEQ ID NO: 808)
PSAT1 (SEQ ID NO: 25)	AATTGTTTTGATTTGGTT (SEQ ID NO: 809)
PSAT1 (SEQ ID NO: 25)	TAATGGGGCGTCGATT (SEQ ID NO: 810)
PSAT1 (SEQ ID NO: 25)	TTAATGGGGTGTTGATT (SEQ ID NO: 811)
PSAT1 (SEQ ID NO: 25)	TATCGTAGCGGTTAGG (SEQ ID NO: 812)
PSAT1 (SEQ ID NO: 25)	TATTGTAGTGGTTAGGAA (SEQ ID NO: 813)
PSAT1 (SEQ ID NO: 25)	AGGAACGTTAGTCGTT (SEQ ID NO: 814)
PSAT1	TAGGAATGTTAGTTGTTT

Gene	Oligo:
(SEQ ID NO: 25)	(SEQ ID NO: 815)
PSAT1 (SEQ ID NO: 25)	GGTCGTCGTATTATGGA (SEQ ID NO: 816)
PSAT1 (SEQ ID NO: 25)	TGGTTGTTGTATTATGGA (SEQ ID NO: 817)
CGA (SEQ ID NO: 26)	ATATTTATTTTCGGAAATTT (SEQ ID NO: 836)
CGA (SEQ ID NO: 26)	TTATTTTGGAAATTTATAGT (SEQ ID NO: 837)
CGA (SEQ ID NO: 26)	TGATTTTGTCGTTATTATT (SEQ ID NO: 838)
CGA (SEQ ID NO: 26)	TTGATTTTGTTGTTATTATT (SEQ ID NO: 839)
CGA (SEQ ID NO: 26)	TAAATTGACGTTATGGTA (SEQ ID NO: 840)
CGA (SEQ ID NO: 26)	AAATTGATGTTATGGTAAA (SEQ ID NO: 841)
CGA (SEQ ID NO: 26)	AATTGACGTTATGGTAAT (SEQ ID NO: 842)
CGA (SEQ ID NO: 26)	TAAAAATTGATGTTATGGT (SEQ ID NO: 843)
COX7A2L (SEQ ID NO: 28)	TTGTTCTGAAGATCGTT (SEQ ID NO: 1078)
COX7A2L (SEQ ID NO: 28)	GTTGTTTGAAGATTGTTT (SEQ ID NO: 1079)

Gene	Oligo:
COX7A2L (SEQ ID NO: 28)	TAGCGTAAGGATTCGGT (SEQ ID NO: 1080)
COX7A2L (SEQ ID NO: 28)	TTAGTGTAAGGATTTGGT (SEQ ID NO: 1081)
COX7A2L (SEQ ID NO: 28)	AGAGTTCGGTTTTTCGTA (SEQ ID NO: 1082)
COX7A2L (SEQ ID NO: 28)	AGAGTTTGGTTTTTTGTA (SEQ ID NO: 1083)
COX7A2L (SEQ ID NO: 28)	ATTCGTATTTGCGGGTTA (SEQ ID NO: 1084)
COX7A2L (SEQ ID NO: 28)	ATTTGTATTTGTGGGTTA (SEQ ID NO: 1085)
ESR2 (SEQ ID NO: 29)	ATTTTCGAGGATTACGTT (SEQ ID NO: 936)
ESR2 (SEQ ID NO: 29)	ATTTTGAGGATTATGTTTT (SEQ ID NO: 937)
ESR2 (SEQ ID NO: 29)	AGATGGCGTTTTTCGTA (SEQ ID NO: 938)
ESR2 (SEQ ID NO: 29)	TAGATGGTGTTTTTTGTA (SEQ ID NO: 939)
ESR2 (SEQ ID NO: 29)	ATTTTCGAATCGATTTTT (SEQ ID NO: 940)
ESR2 (SEQ ID NO: 29)	GGAGTATTTTTGAATTGAT (SEQ ID NO: 941)
ESR2 (SEQ ID NO: 29)	AGTTCGACGGTTTGTAG (SEQ ID NO: 942)

Gene	Oligo:
29)	
ESR2 (SEQ ID NO: 29)	AGGGAGTTTGATGGTT (SEQ ID NO: 943)
ESR2 (SEQ ID NO: 29)	AGTTTACGTGATCGAG (SEQ ID NO: 944)
ESR2 (SEQ ID NO: 29)	AGTTTATGTGATTGAGTT (SEQ ID NO: 945)
VTN (SEQ ID NO: 31)	GGTGGTATCGATTGAT (SEQ ID NO: 1034)
VTN (SEQ ID NO: 31)	TGGTGGTATTGATTGAT (SEQ ID NO: 1035)
VTN (SEQ ID NO: 31)	TAGTGATTCGCGGGGA (SEQ ID NO: 1038)
VTN (SEQ ID NO: 31)	TAGTGATTTGTGGGGA (SEQ ID NO: 1039)
VTN (SEQ ID NO: 31)	TTATGTCGGAGGATGA (SEQ ID NO: 1040)
VTN (SEQ ID NO: 31)	ATTATGTTGGAGGATGA (SEQ ID NO: 1041)
VTN (SEQ ID NO: 31)	ATACGGTTTATGACGAT (SEQ ID NO: 1042)
VTN (SEQ ID NO: 31)	ATATGGTTTATGATGATGG (SEQ ID NO: 1043)
PCAF (SEQ ID NO: 33)	GAGCGGTAGGTGTCGAA (SEQ ID NO: 978)
PCAF	GAGTGGTAGGTGTTGAA

Gene	Oligo:
(SEQ ID NO: 33)	(SEQ ID NO: 979)
PCAF (SEQ ID NO: 33)	TAAGATTTTCGCGGGTA (SEQ ID NO: 980)
PCAF (SEQ ID NO: 33)	TGTAAGATTTTGTGGGTA (SEQ ID NO: 981)
PCAF (SEQ ID NO: 33)	AGTTCGTAGTTTCGAG (SEQ ID NO: 982)
PCAF (SEQ ID NO: 33)	GTTTGTAGTTTTGAGGA (SEQ ID NO: 983)
PCAF (SEQ ID NO: 33)	TAGGGCGCGGAGTAGA (SEQ ID NO: 984)
PCAF (SEQ ID NO: 33)	TAGGGTGTGGAGTAGA (SEQ ID NO: 985)
PRKCD (SEQ ID NO: 34)	ATTTATTTTTCGTTGTAGG (SEQ ID NO: 772)
PRKCD (SEQ ID NO: 34)	TATTTATTTTGTGTAGG (SEQ ID NO: 773)
PRKCD (SEQ ID NO: 34)	TTTCGGAAACGGGAAT (SEQ ID NO: 774)
PRKCD (SEQ ID NO: 34)	TAGTTTTGGAAATGGGA (SEQ ID NO: 775)
PRKCD (SEQ ID NO: 34)	GGACGGAGTTATCGGT (SEQ ID NO: 776)
PRKCD (SEQ ID NO: 34)	GGATGGAGTTATTGGTA (SEQ ID NO: 777)

Gene	Oligo:
PRKCD (SEQ ID NO: 34)	GTTTAGCGGAGGGATA (SEQ ID NO: 778)
PRKCD (SEQ ID NO: 34)	TGTTTAGTGGAGGGAT (SEQ ID NO: 779)
ESR1 (exon8) (SEQ ID NO: 61)	TTGTTACGGTTTGAGAG (SEQ ID NO: 780)
ESR1 (exon8) (SEQ ID NO: 61)	TTGTTATGGTTTGAGAGT (SEQ ID NO: 781)
ESR1 (exon8) (SEQ ID NO: 61)	TTTGTTATAGTTTGAGAGT (SEQ ID NO: 782)
ESR1 (exon8) (SEQ ID NO: 61)	TTTGTTACGGTTTGAG (SEQ ID NO: 783)
ESR1 (exon8) (SEQ ID NO: 61)	TTTGTTATGGTTTGAGA (SEQ ID NO: 784)
ESR1 (exon8) (SEQ ID NO: 61)	TTTGTTATAGTTTGAGAG (SEQ ID NO: 785)

10 -02- 2004

I/We claim:

1. A method for characterising a cell proliferative disorder of the breast tissues and/or a metastases thereof and/or predicting the disease free survival and/or response of a subject with said disorder to a treatment comprising one or more treatment which target the estrogen receptor pathway or are involved in estrogen metabolism, production or secretion, said method comprising

- a) obtaining a biological sample from the subject
- b) determining the methylation status of one or more CpG positions within at least one target nucleic acid comprising one or a combination of the genes taken from the group consisting of EGR4, APC, CDKN2A, CSPG2, ERBB2, STMN1, STK11, CA9, PAX6, SFN, S100A2, TFF1, TGFBR2, TP53, TP73, PLAUI, TMEFF2, ESR1, SYK, HSPB1, RASSF1, TES, PITX2, GRIN2D, PSAT1, CGA, CYP2D6, COX7A2L, ESR2, PLAUI, VTN, SULT1A1, PCAF, PRKCD, ONECUT2, BCL6, WBP11, (MX1), N.N., APP, ORC4L, NETO1, TBC1D3, GRB7, CYP2D6, CDK6, (Chr. 1p13.2), (Chr. 17q25.1), ABCA8, (Chr. 12q14.3), (Chr. 8q12.1), MARK2, ELK1, Q8WUT3, CGB, BSG, BCKDK, SOX8, DAG1, SEMA4B, ESR1 (exon8) and/or their regulatory regions by contacting said target nucleic acid with one or more agents that convert cytosine bases that are unmethylated at the 5'-position thereof to a base that is detectably dissimilar to cytosine in terms of hybridization properties
- c) determining therefrom the prognosis of said subject, characteristics of said cell proliferative disorder, disease free survival and/or probability of response of said subject to said treatment

2. The method according to claims 1 further comprising

- d) determining a suitable treatment regimen for the subject

3. The method according to claims 1 and 2 wherein said suitable treatment regimen comprises one or more therapies selected from the group consisting of

chemotherapy, radiotherapy, surgery, biological therapy, immunotherapy, antibodies, molecularly targeted drugs, estrogen receptor modulators, estrogen receptor down-regulators, aromatase inhibitors, ovarian ablation, LHRH analogues and other centrally acting drugs influencing estrogen production.

4. A method according to Claims 1 to 3, wherein said treatment is an adjuvant treatment and said genes are selected from the group consisting of ERBB2, STMN1, TFF1, TMEFF2, ESR1, HSPB1, PITX2, COX7A2L, PLAU, VTN, PCAF, ONECUT2, BCL6, WBP11, TBC1D3, GRB7, CDK6, (Chr. 1p13.2), ABCA8 and (Chr. 8q12.1)

5. A method according to Claims 1 to 3, wherein said treatment is an adjuvant treatment and said target nucleic acid(s) are selected from the group consisting of SEQ ID NO: 5, 6, 12, 17, 18, 20, 23, 28, 16, 31, 33, 35, 36, 37, 43, 44, 46, 47, 49 and 51.

6. A method according to Claim 1 to 3, wherein said disorder is a metastatic disease and said genes are selected from the group consisting of APC, CSPG2, ERBB2, STK11, S100A2, TFF1, TGFBR2, TP53, TMEFF2, SYK, HSPB1, RASSF1, PSAT1, CGA, ESR2, ONECUT2, WBP11, CYP2D6, CDK6, ELK1, CGB and DAG1

7. A method according to Claims 1 to 3, wherein said disorder is a metastatic disease and said target nucleic acid(s) are selected from the group consisting of SEQ ID NO: 2, 4, 5, 7, 11, 12, 13, 14, 17, 19, 20, 21, 25, 26, 29, 35, 37, 45, 46, 53, 55 and 59.

8. The method as recited in one of the Claims 1 through 7, characterised in that the genomic DNA is obtained from cells or cellular components which contain DNA, sources of DNA comprising, for example, cell lines, histological slides, biopsies, tissue embedded in paraffin or sections thereof, breast tissues, blood, plasma, serum, lymphatic

fluid, lymphatic tissue, duct cells, ductal lavage fluid, nipple aspiration fluid, cerebrospinal fluid, bone marrow and combinations thereof.

9. A method according to Claims 1 to 8, wherein said cell proliferative disorder of the breast tissue is selected from the group consisting of ductal carcinoma *in situ*, invasive ductal carcinoma, invasive lobular carcinoma, lobular carcinoma *in situ*, comedocarcinoma, inflammatory carcinoma, mucinous carcinoma, scirrhous carcinoma, colloid carcinoma, tubular carcinoma, medullary carcinoma, metaplastic carcinoma, and papillary carcinoma and papillary carcinoma *in situ*, undifferentiated or anaplastic carcinoma and Paget's disease of the breast.

10. A method according to Claims 1 to 9, wherein said subjects are estrogen and/or progesterone receptor positive.

11. A method according to claims 1 to 10 wherein b) comprises

- a. converting cytosine bases in the genomic DNA sample which are unmethylated at the 5-position, to uracil or another base which is dissimilar to cytosine in terms of base pairing behaviour;
- b. amplifying at least one fragment of the pretreated genomic DNA, wherein said fragments comprise at least 8 base pairs of one or more sequences selected from the group consisting of SEQ ID NO: 206 to 449 and sequences complementary thereto, and
- c. determining the methylation status of one or more genomic CpG dinucleotides by analysis of the amplificate nucleic acids.

12. The method according to claim 11 wherein ii) is carried out using one or both of MSP and/or HeavyMethyl.

13. The method according to claim 11 wherein iii) is carried out by means of one or more methods taken from the group consisting oligonucleotide hybridisation analysis, Ms SnuPE, sequencing, Real Time detection probes and oligonucleotide array analysis.
14. A nucleic acid molecule consisting essentially of a sequence at least 18 bases in length according to one of the sequences taken from the group consisting of SEQ ID Nos: 206 to 449.
15. An oligomer, in particular an oligonucleotide or peptide nucleic acid (PNA)-oligomer, said oligomer consisting essentially of at least one base sequence having a length of at least 10 nucleotides which hybridises to or is identical to one of the nucleic acid sequences according to SEQ ID NO: 206 to 449.
16. A set of at least two oligonucleotides as recited in claim 15.
17. A kit comprising a bisulfite (= disulfite, hydrogen sulfite) reagent as well as oligonucleotides and/or PNA-oligomers according to one of the Claims 15 or 16.
18. A kit according to claim 17, further comprising standard reagents for performing a methylation assay from the group consisting of MS-SNuPE, MSP, Methyl light, Heavy Methyl, nucleic acid sequencing and combinations thereof.
19. The use of a method according to one of claims 1 through 13, a nucleic acid according to Claim 14, of an oligonucleotide or PNA-oligomer according to Claim 15, of a kit according to Claim 17 or 18 or of a set of oligonucleotides according to claim 16 for the treatment of breast cell proliferative disorders.

Abstract

The present invention relates to modified and genomic sequences, to oligonucleotides and/or PNA-oligomers for detecting the cytosine methylation state of genomic DNA, as well as to a method for predicting the disease free survival and/or response of a subject with a cell proliferative disorder of the breast tissues, to endocrine treatment.



Figure 1

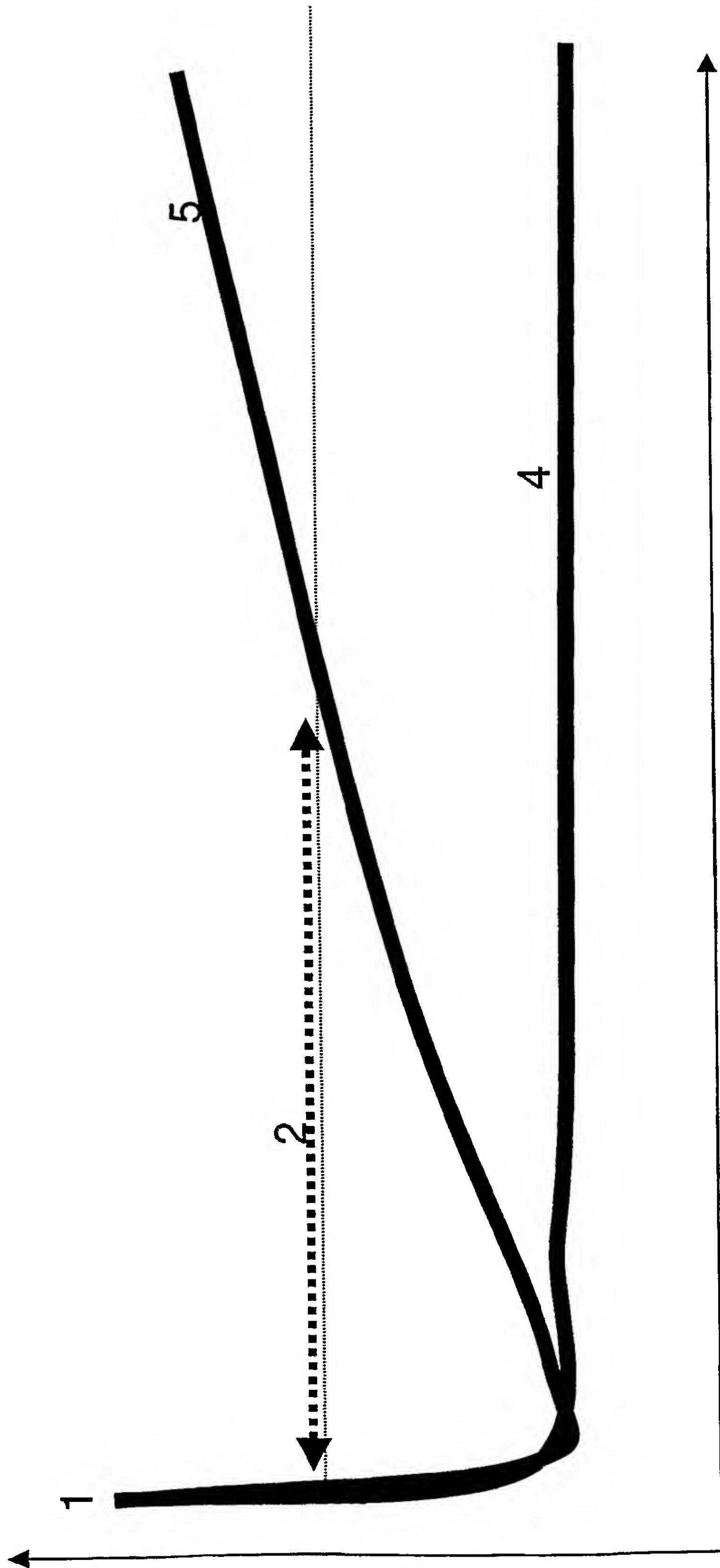
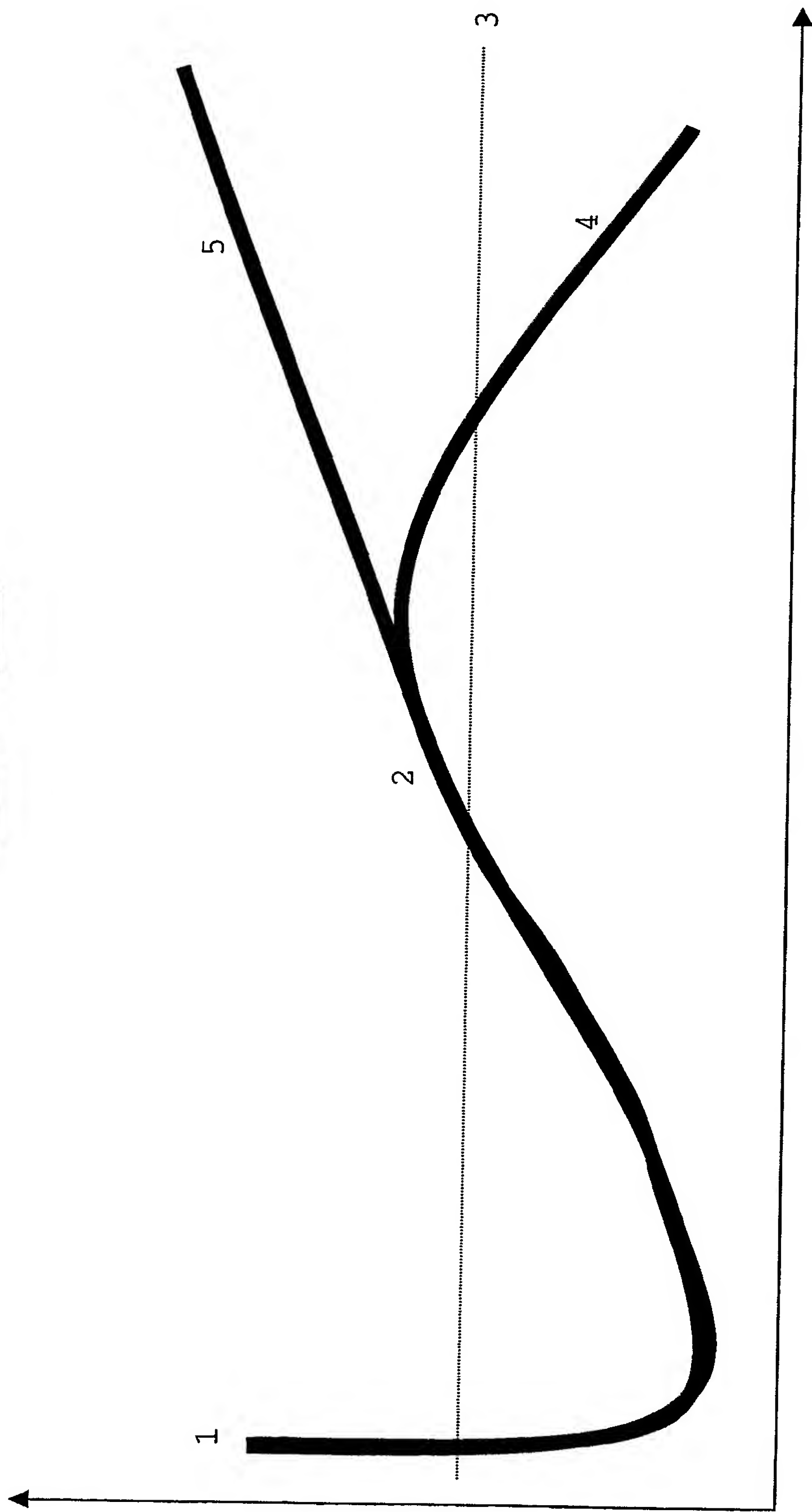


Figure 2



Marker ABCA8 (N= 278)

FIGURE 3

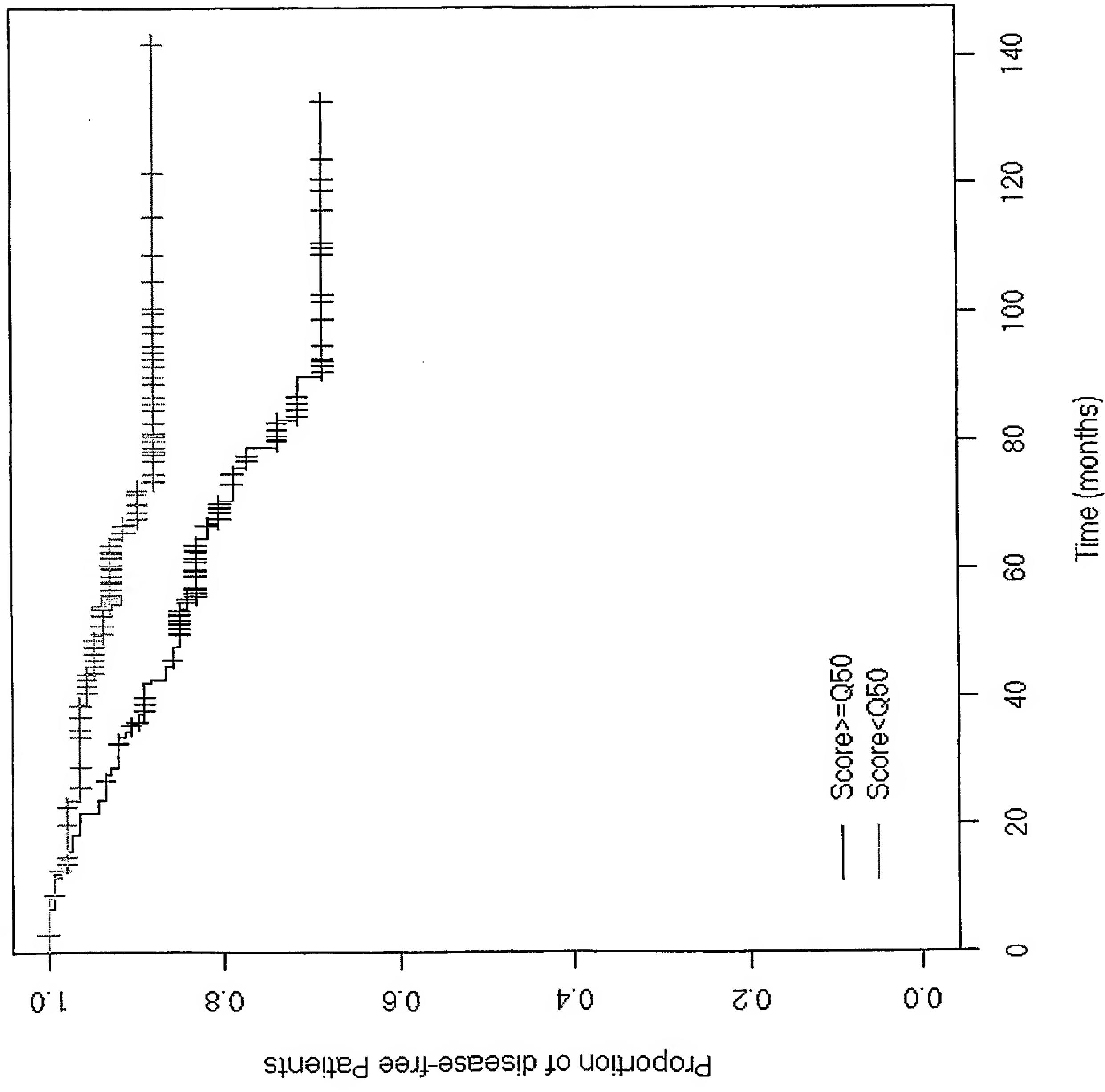
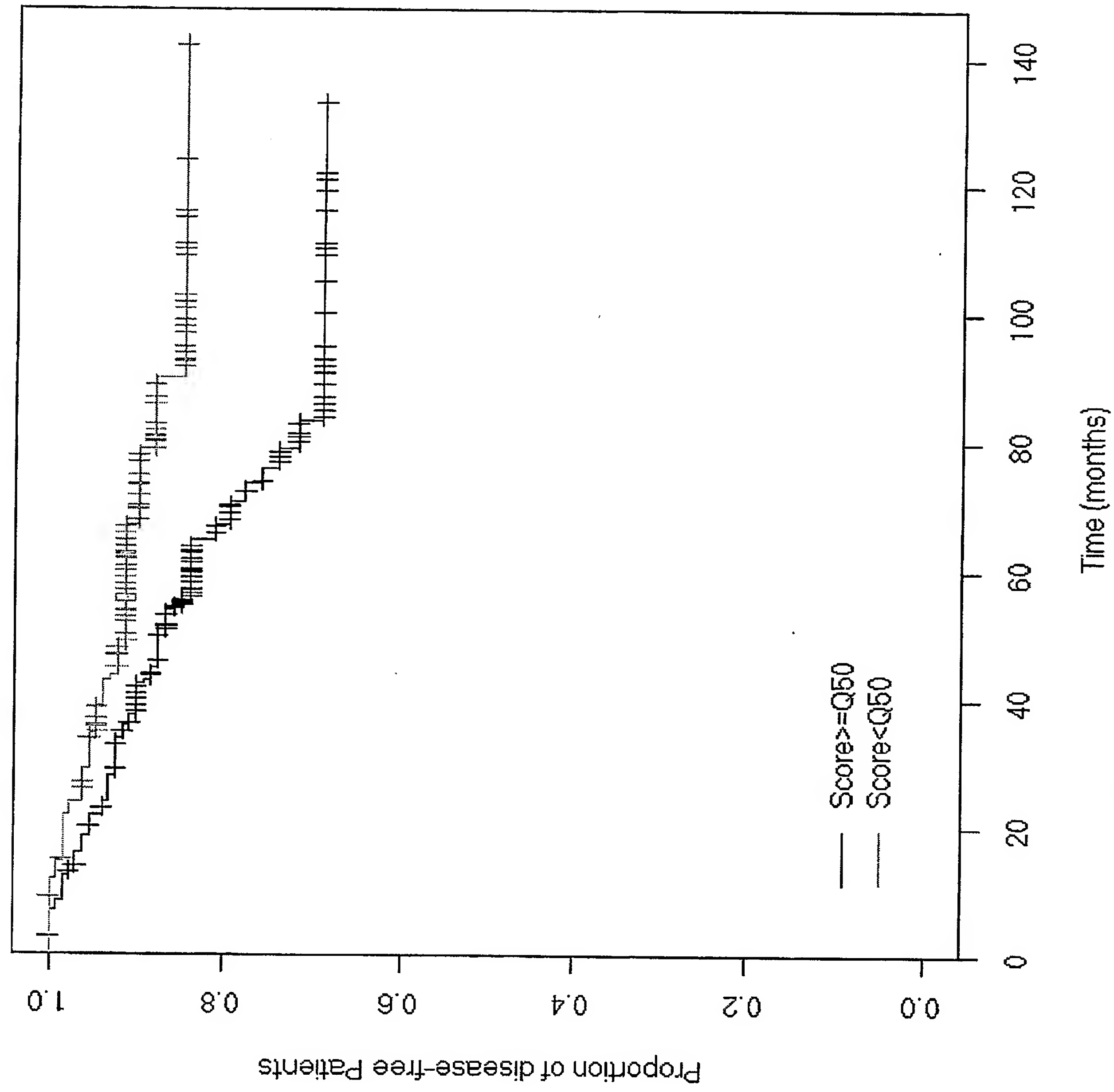


FIGURE 4

Marker BCL6 (N= 278)



Marker CDK6 (N= 278)

FIGURE 5

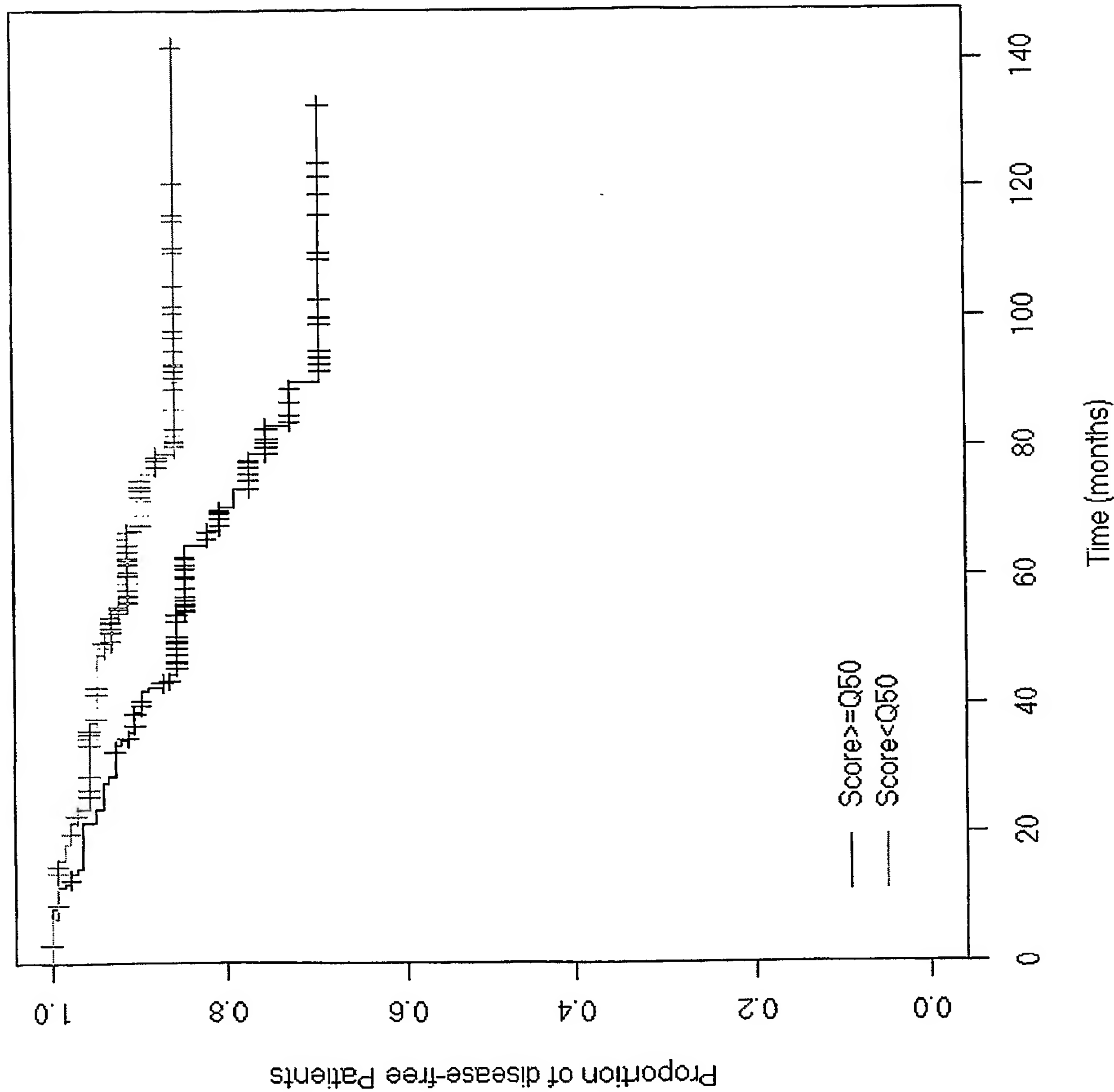


FIGURE 6 **Marker PITX2 (N= 278)**

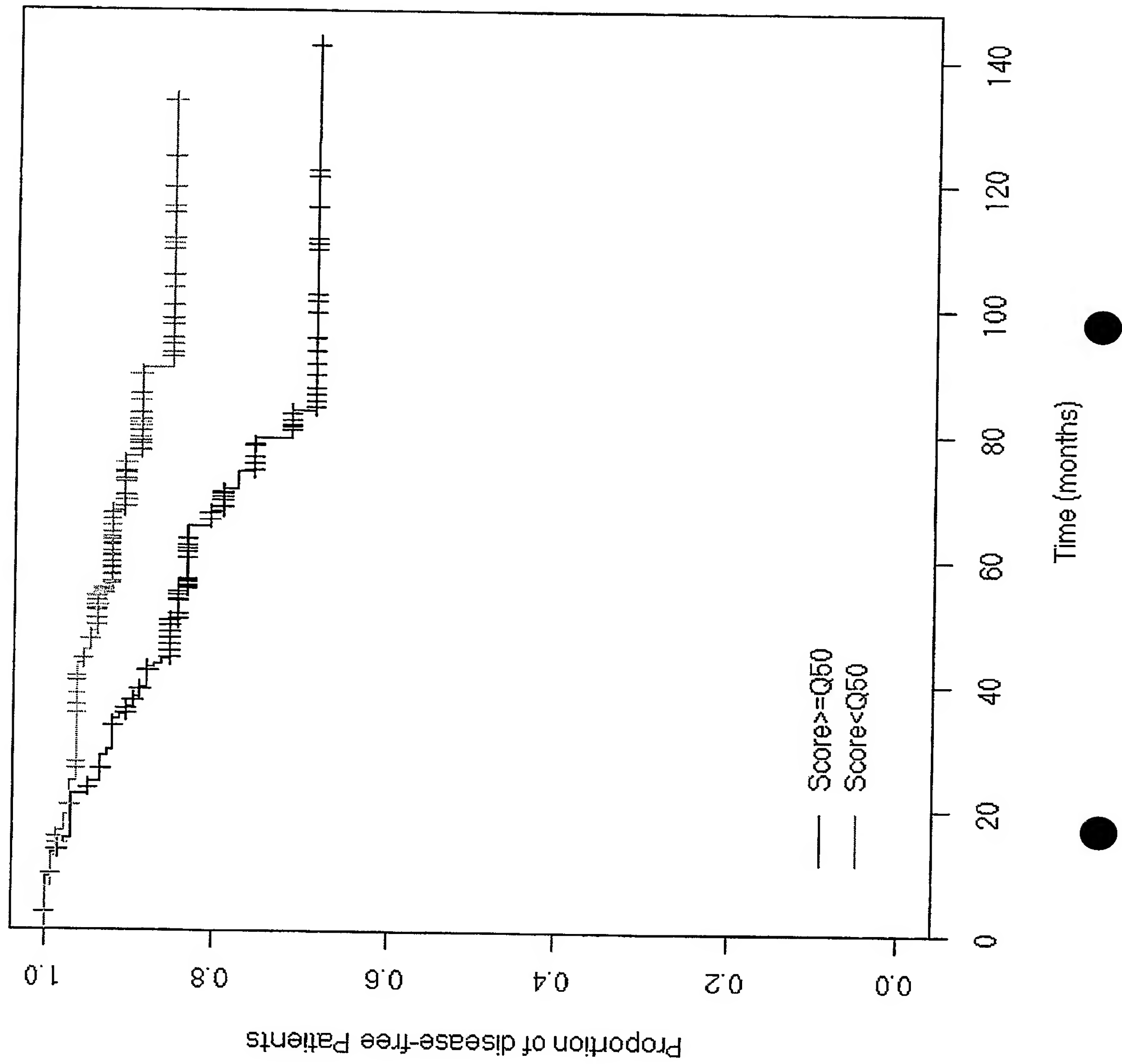


FIGURE 7 Marker STMN1 (N= 278)

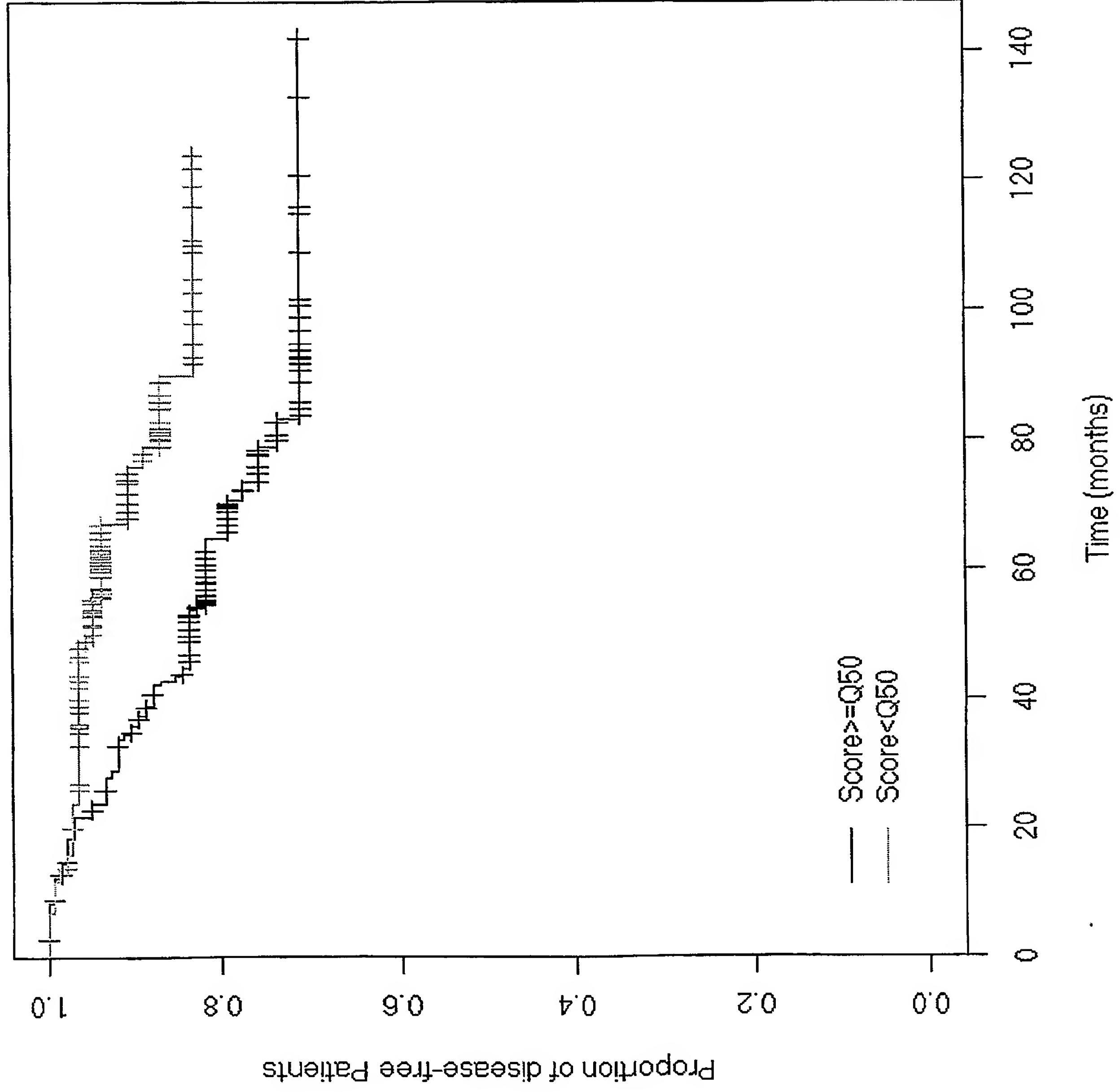
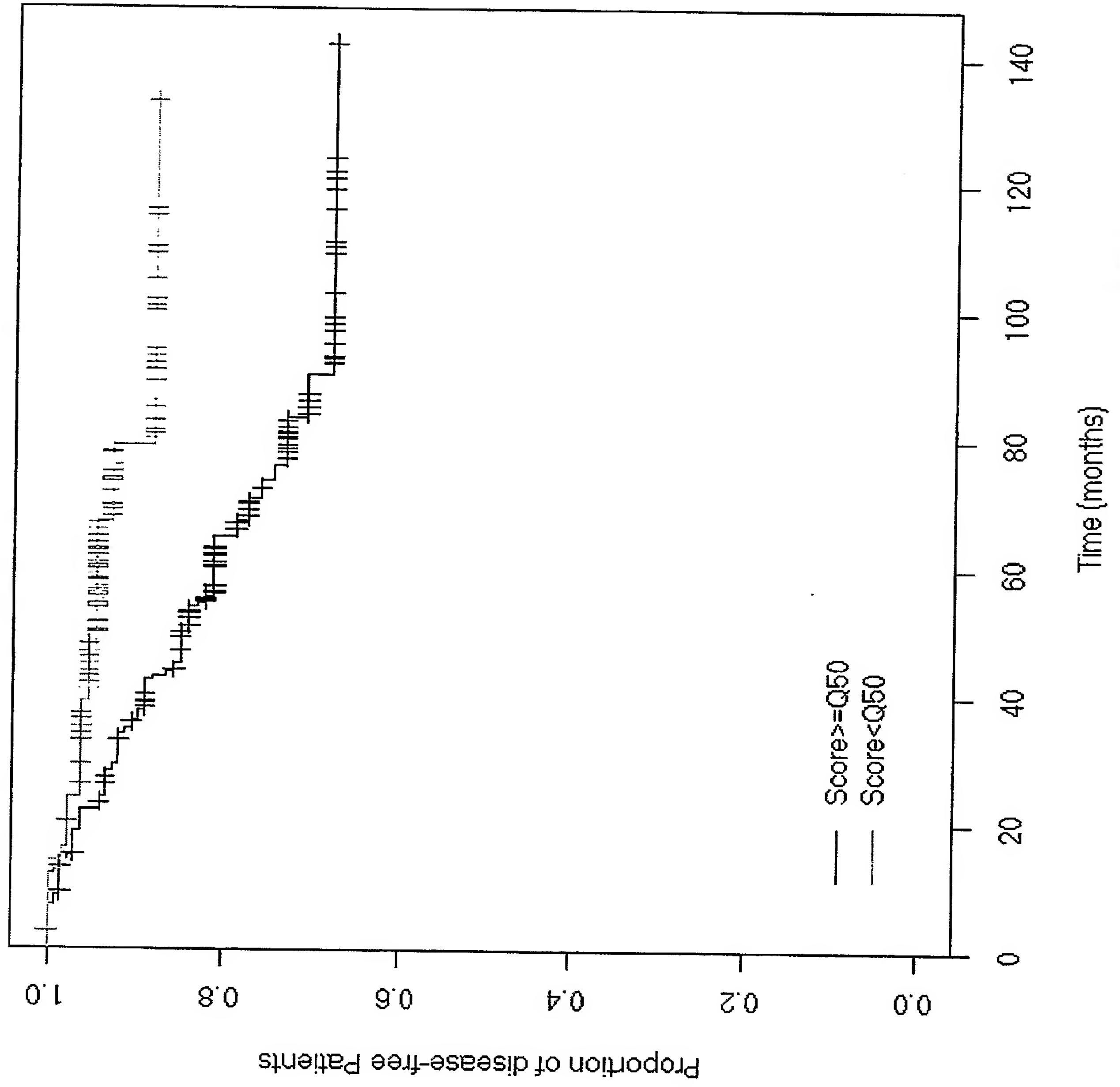


FIGURE 8

Marker TBC1D3 (N= 278)



Marker VTN (N= 278)

FIGURE 9

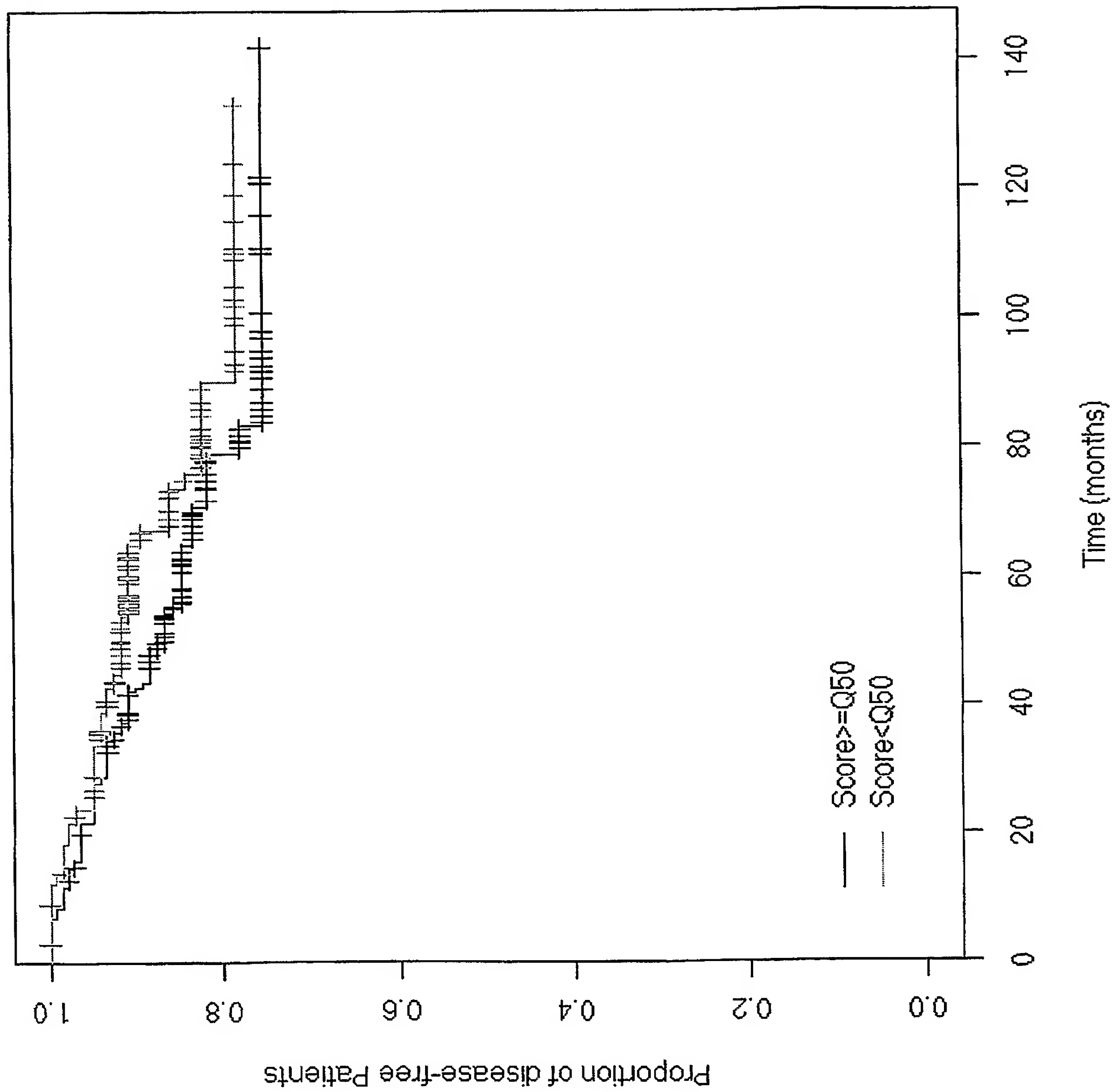


FIGURE 10

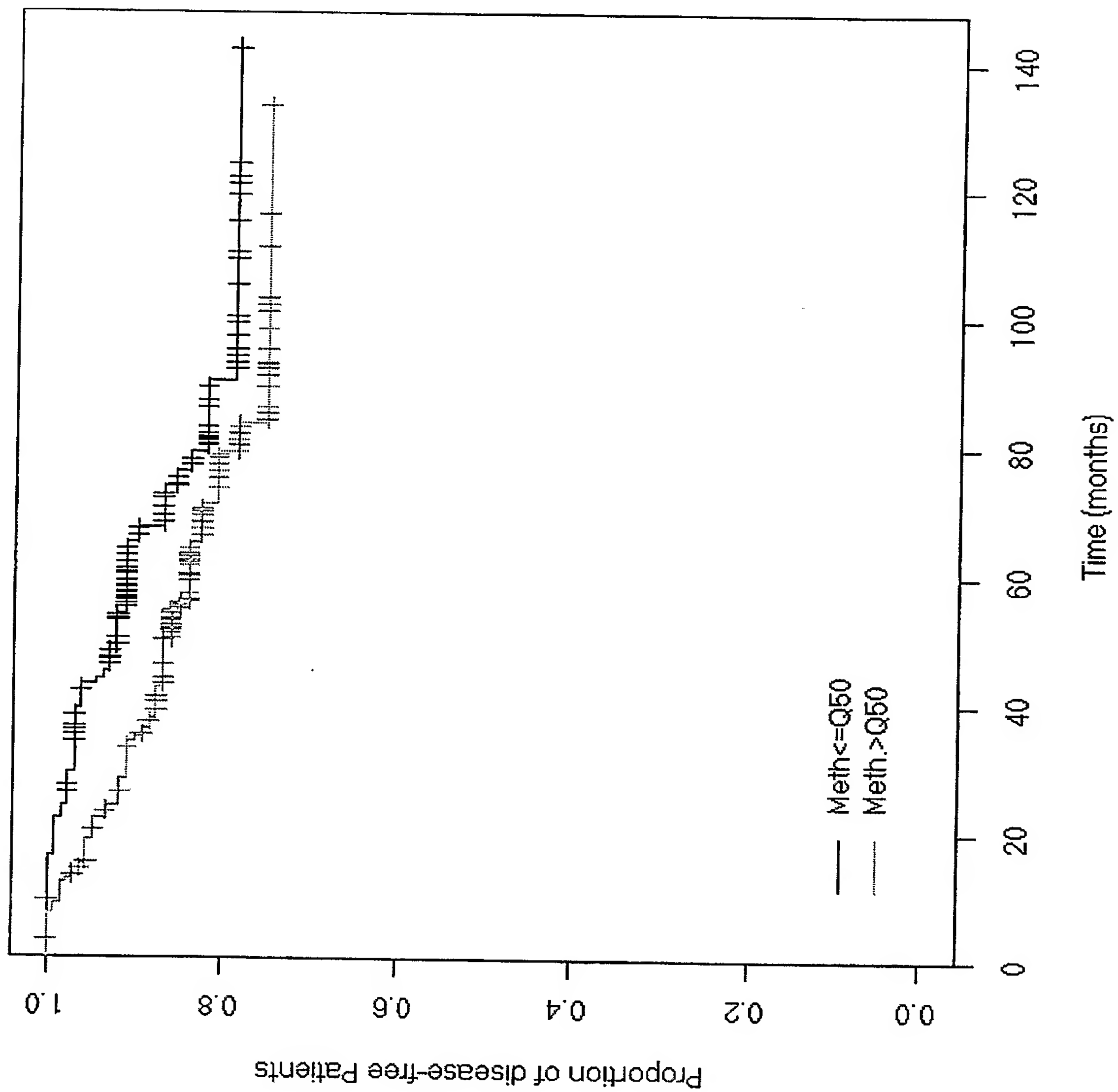


FIGURE 11

SEQ ID NO: 888

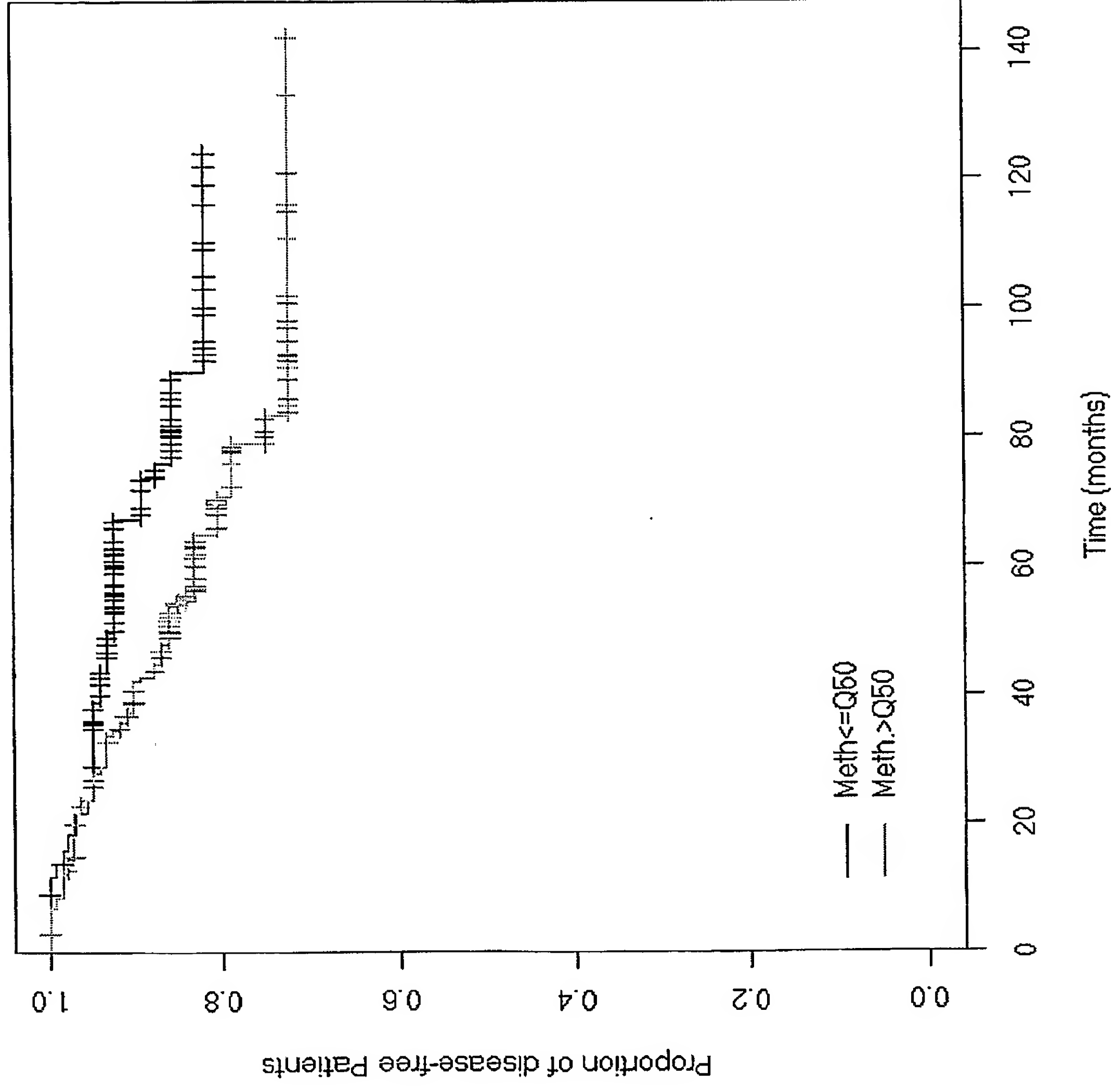


FIGURE 12

SEQ ID NO: 1008

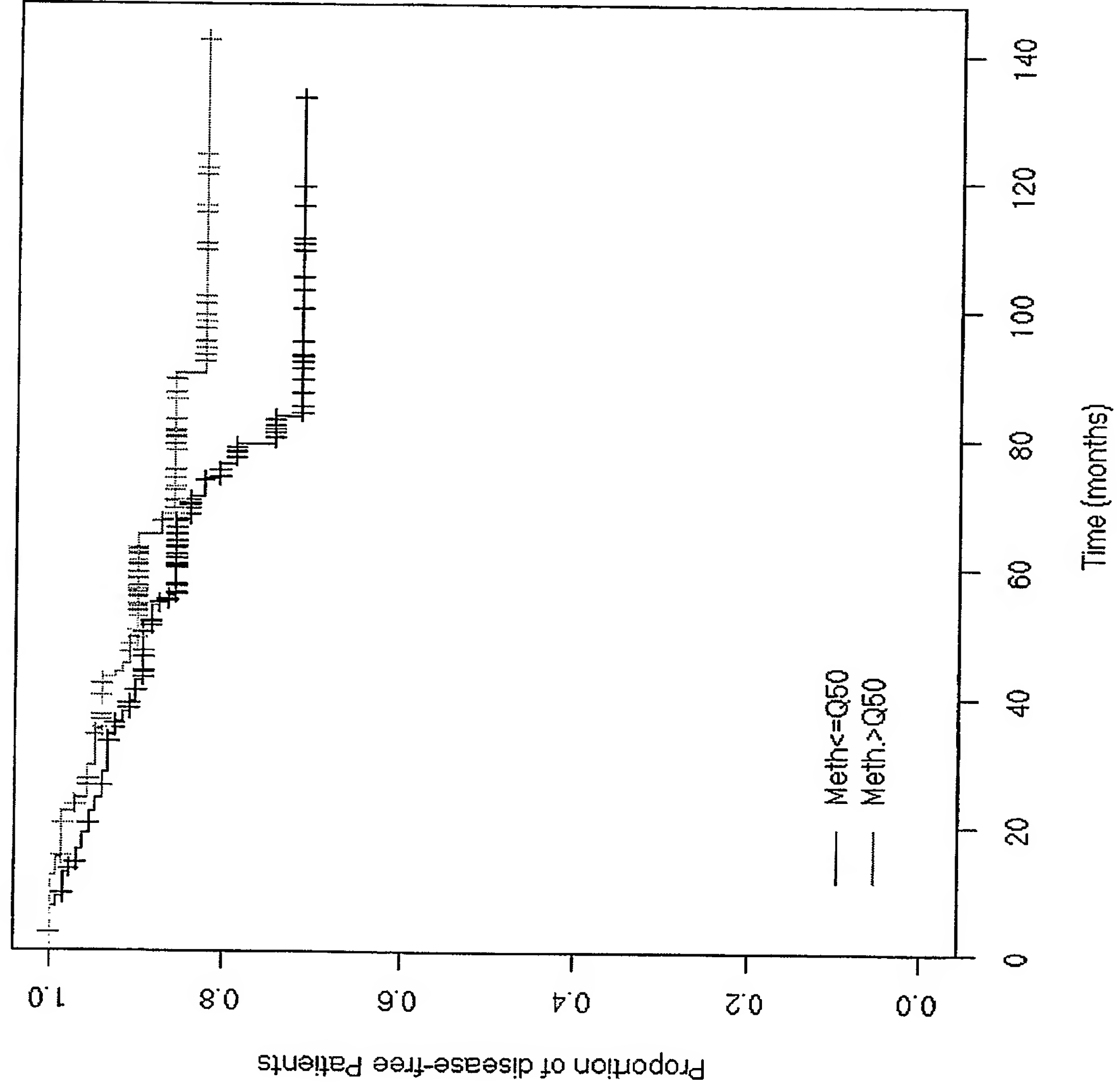


FIGURE 13

SEQ ID NO:794

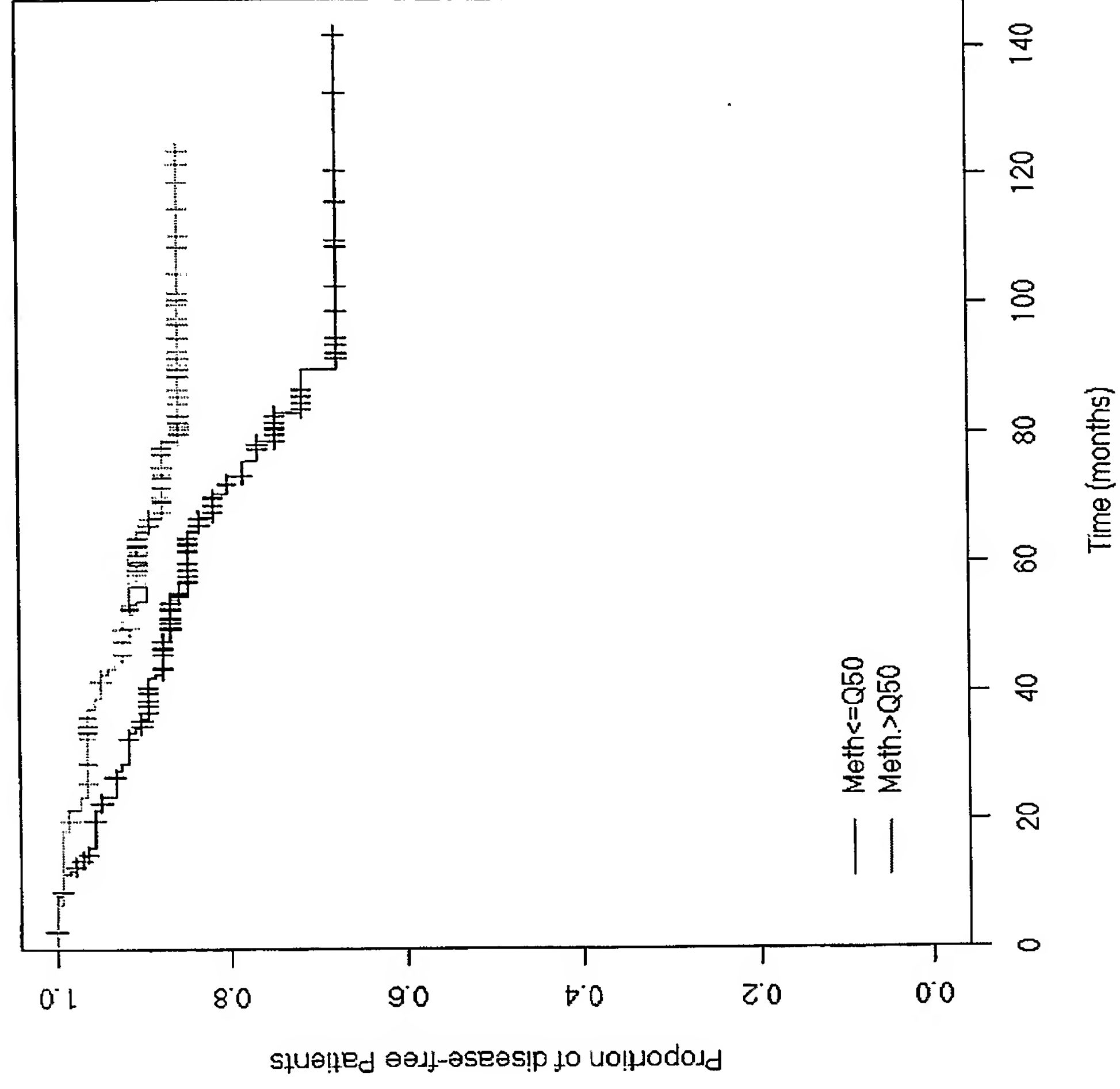


FIGURE 14 (SEQ ID NO: 980)

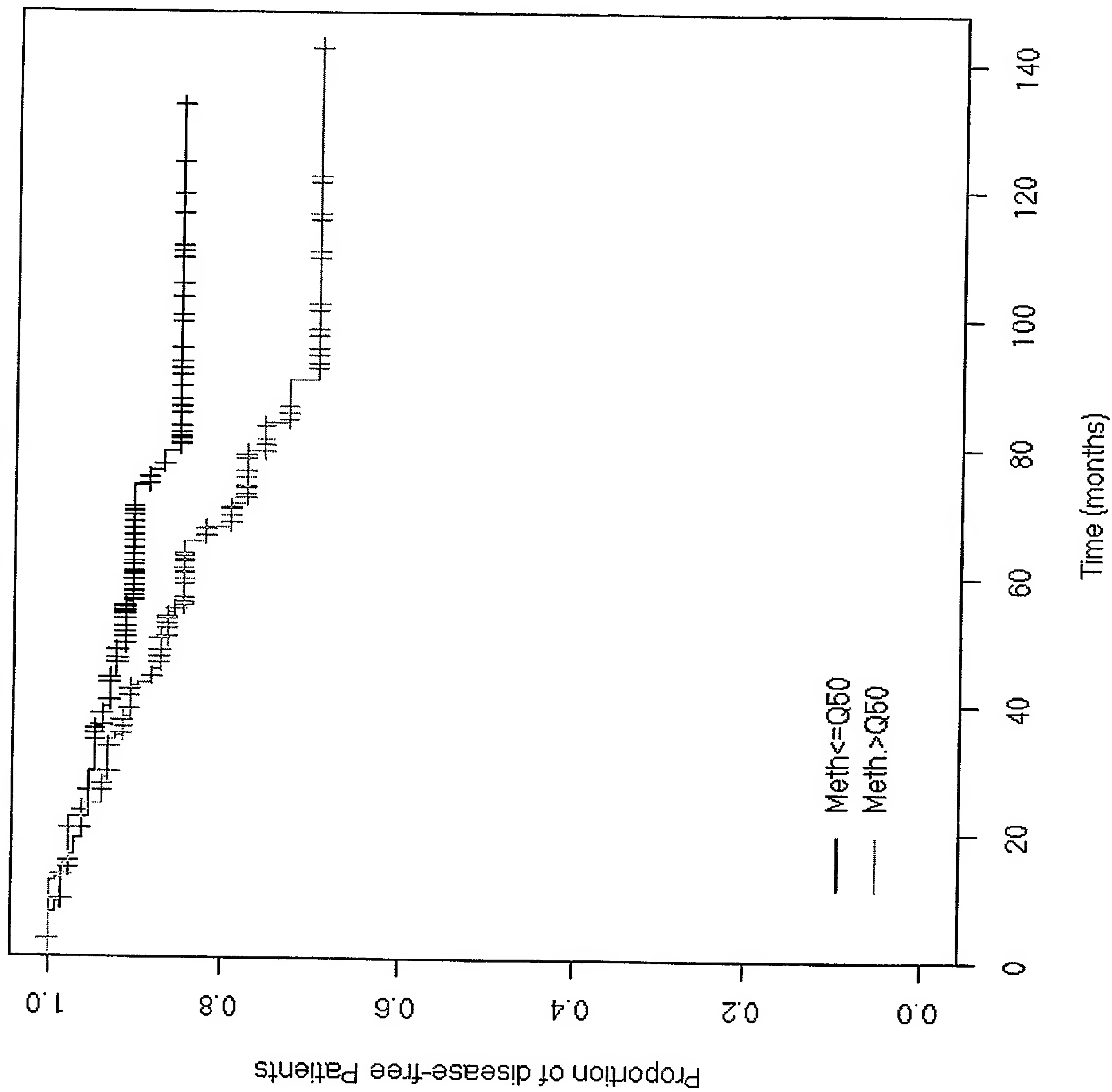


Figure 15 SEQ ID NO: 914

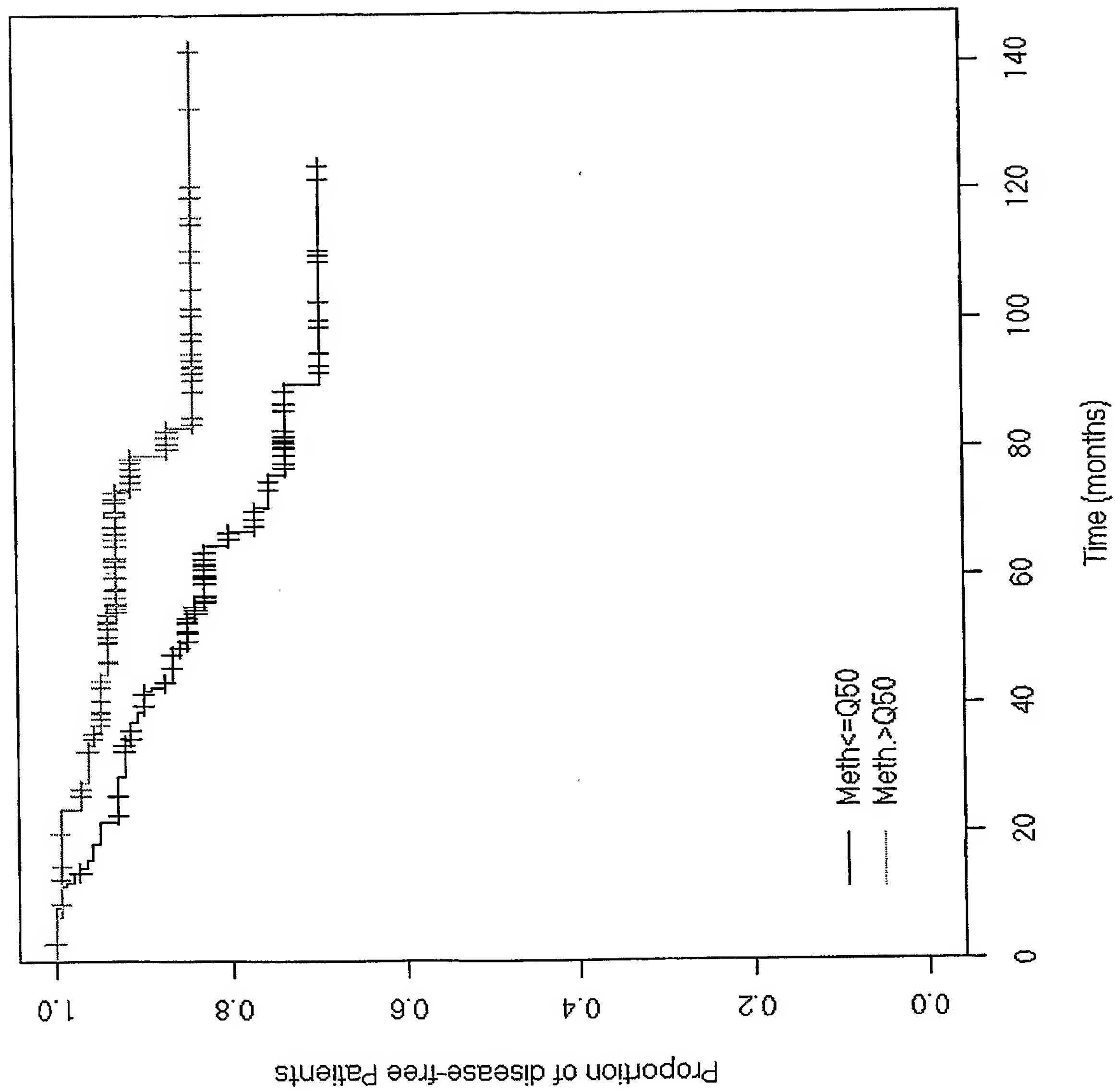


Figure 16 SEQ ID NO: 806

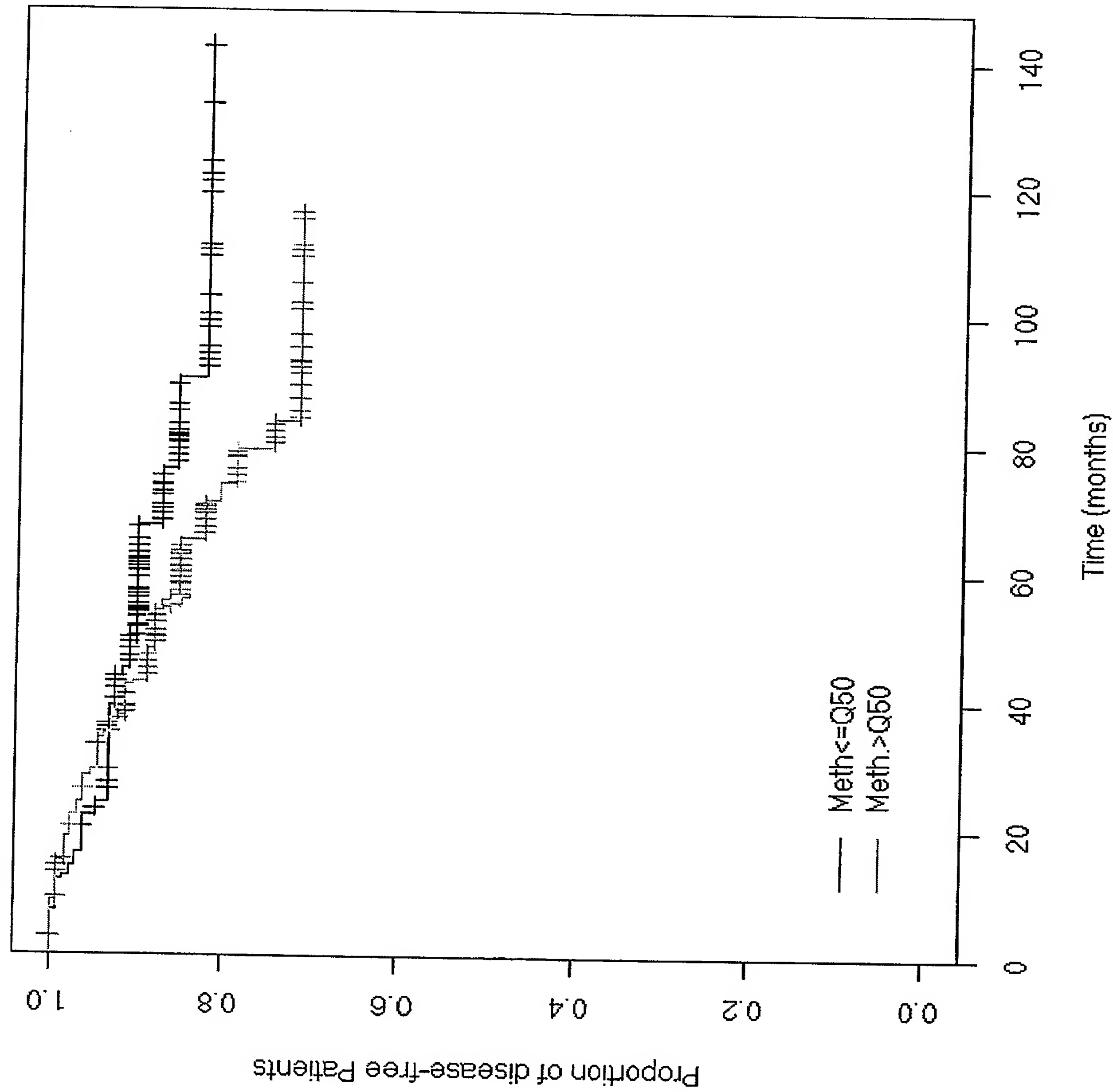


Figure 17 SEQ ID NO: 966

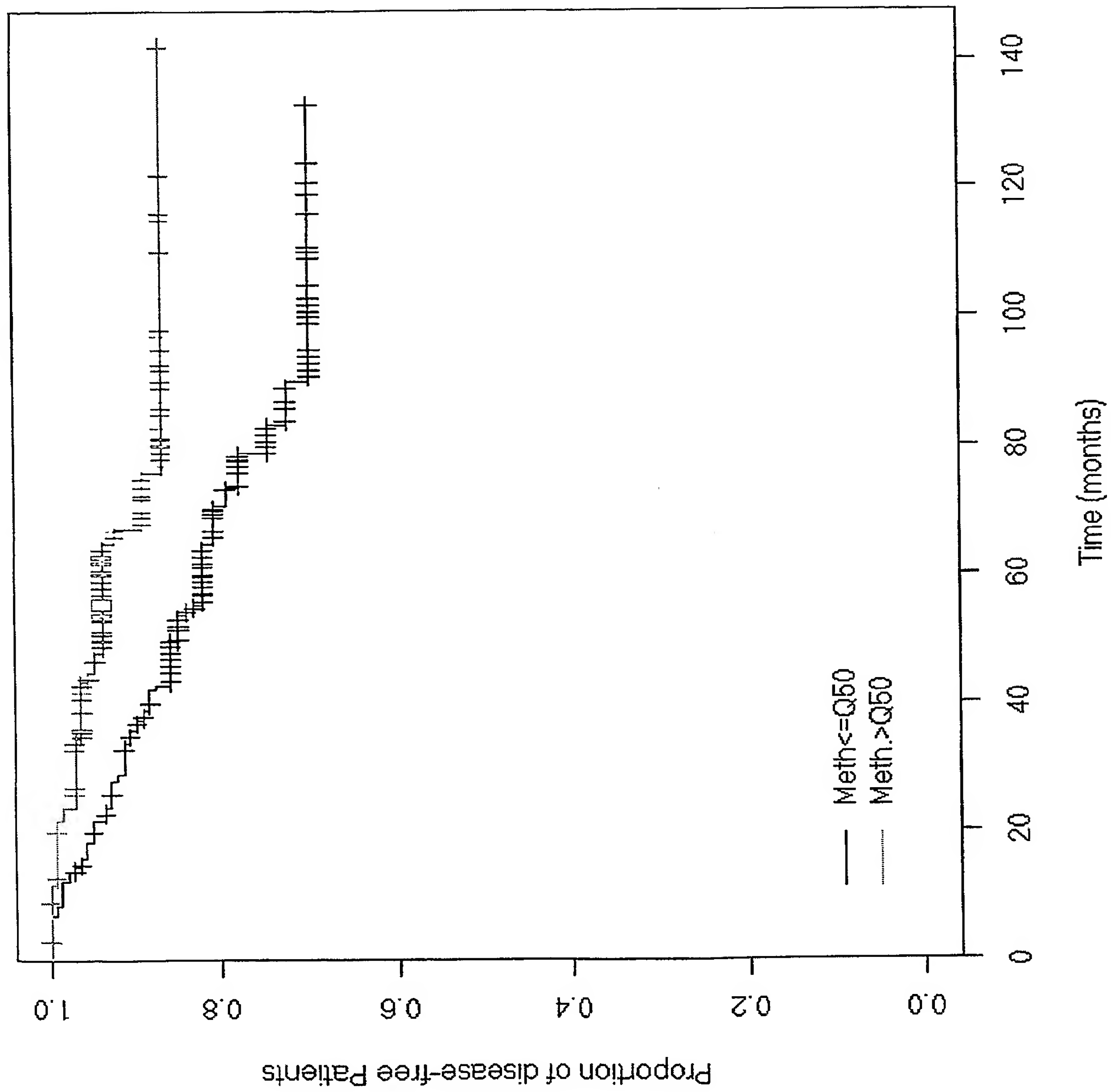


Figure 18 SEQ ID NO: 804

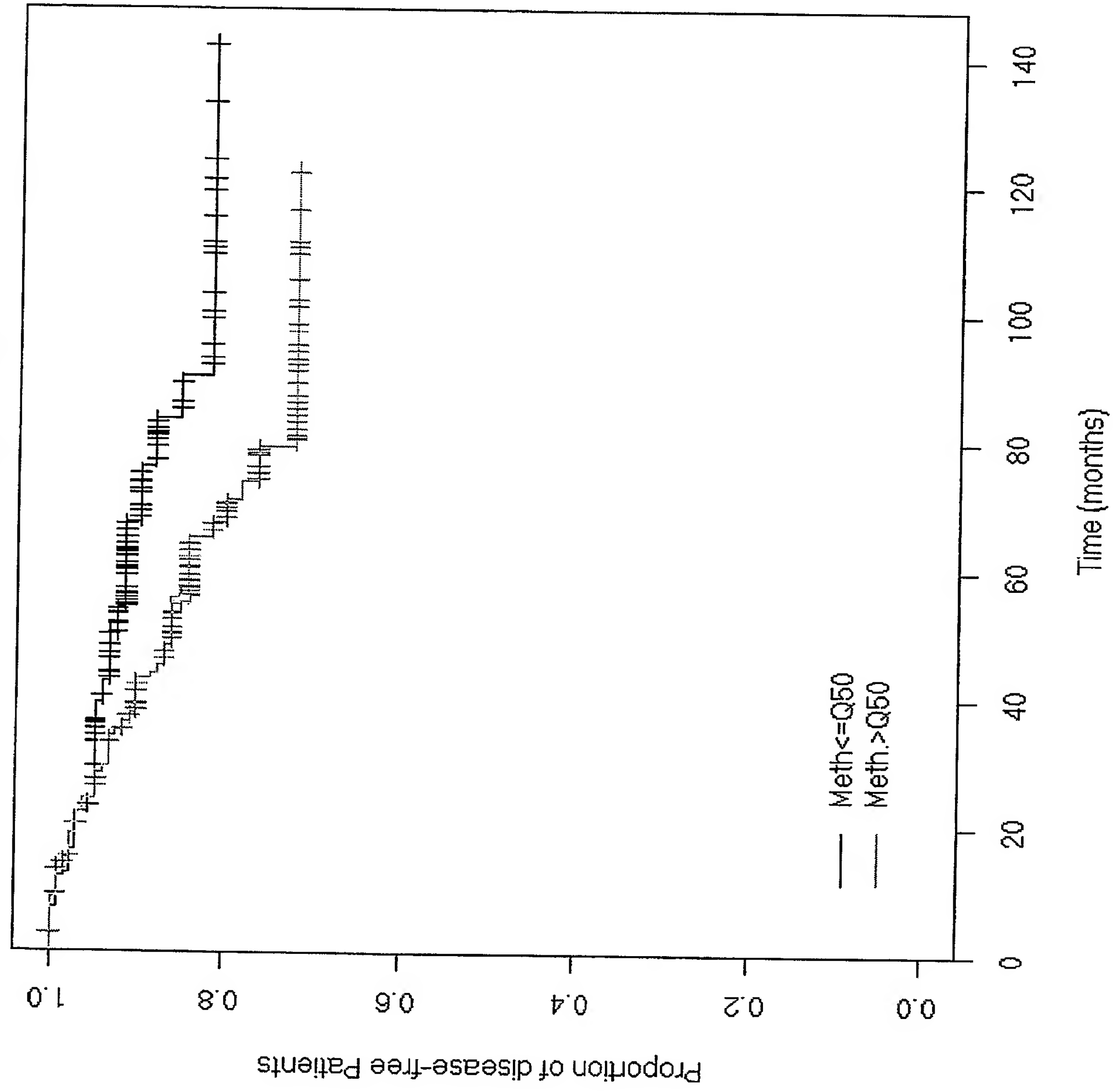


Figure 19 SEQ ID NO 1076

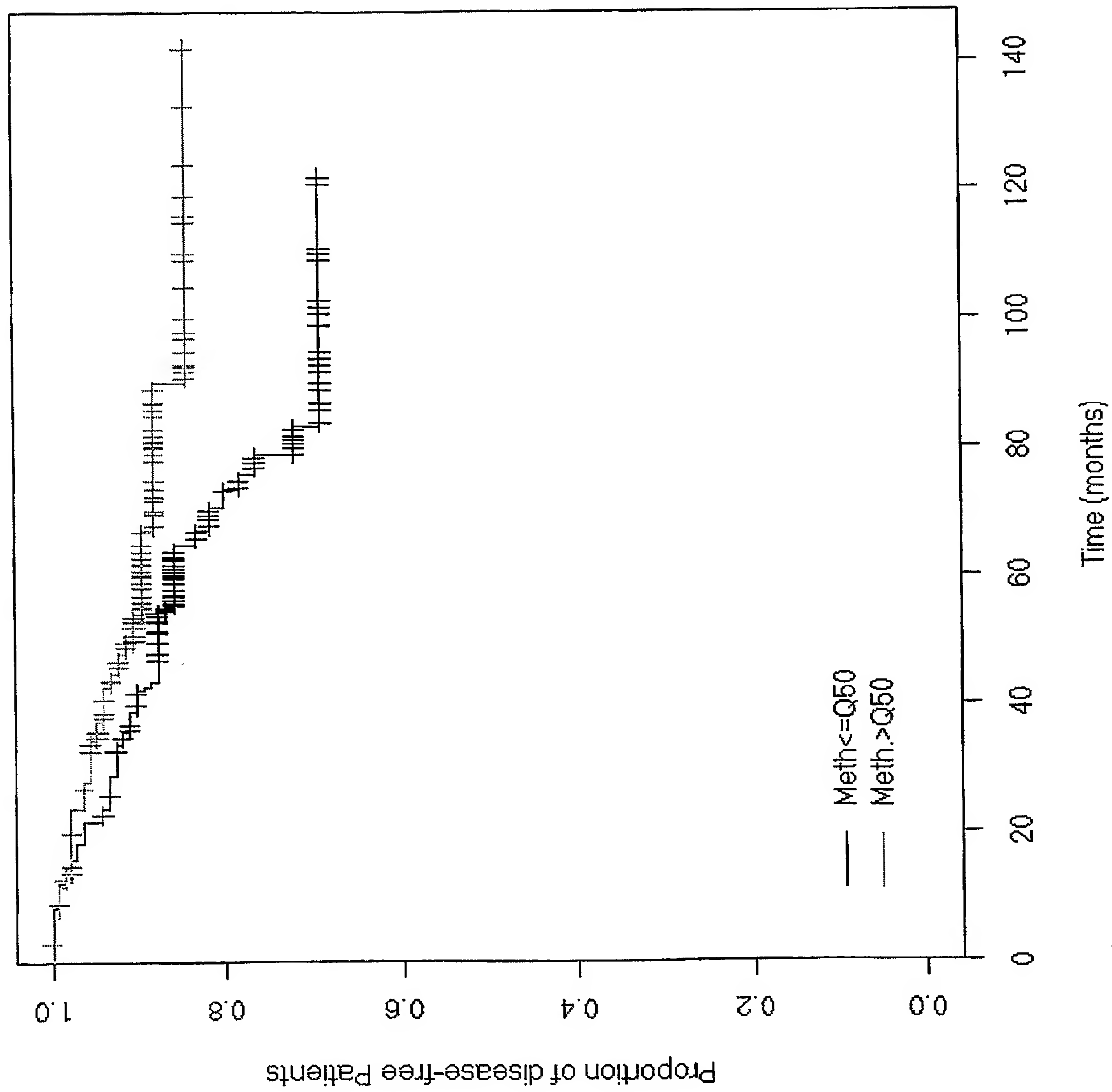


Figure 20 SEQ ID NO: 618

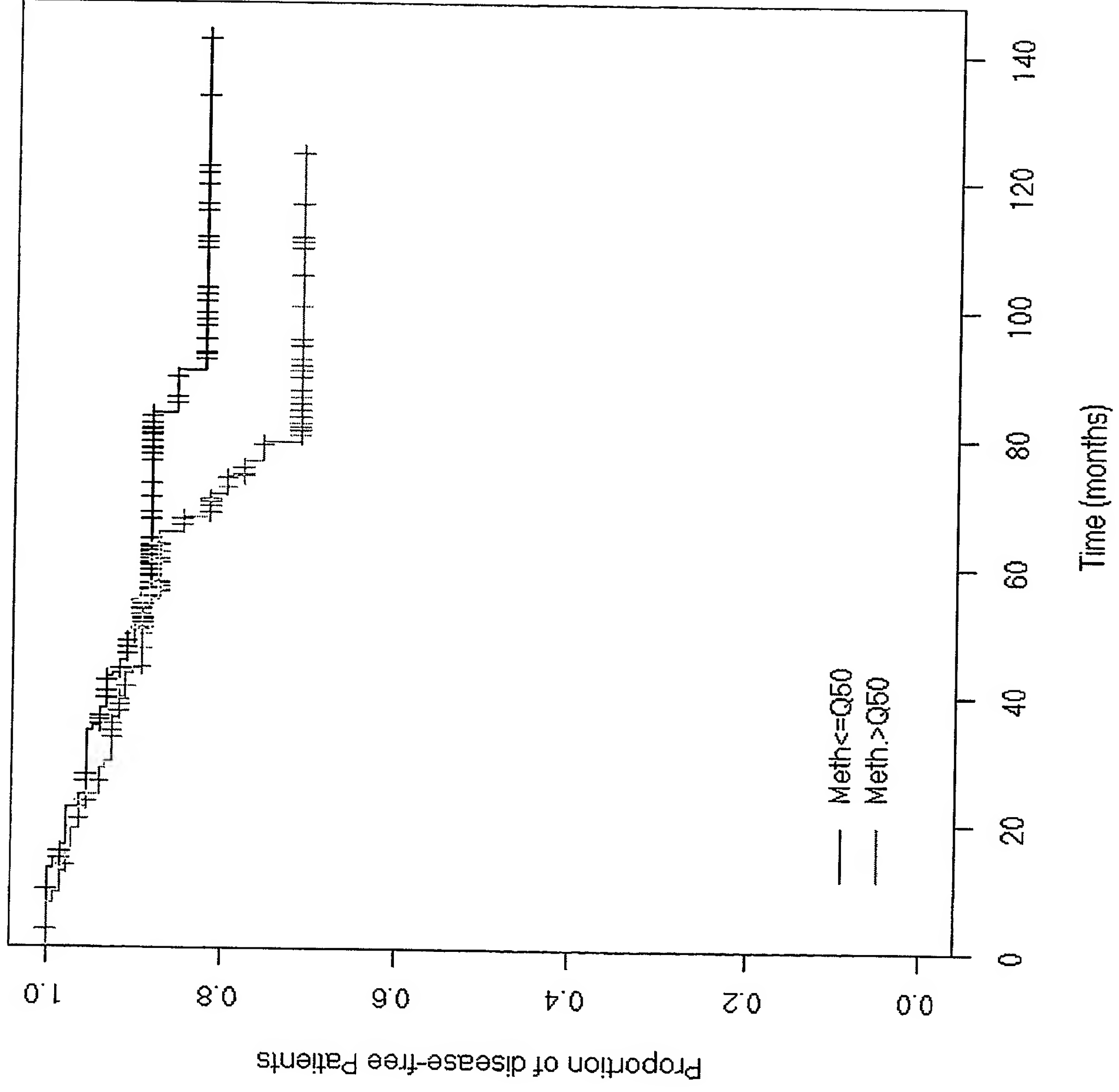


Figure 21 SEQ ID NO: 1054

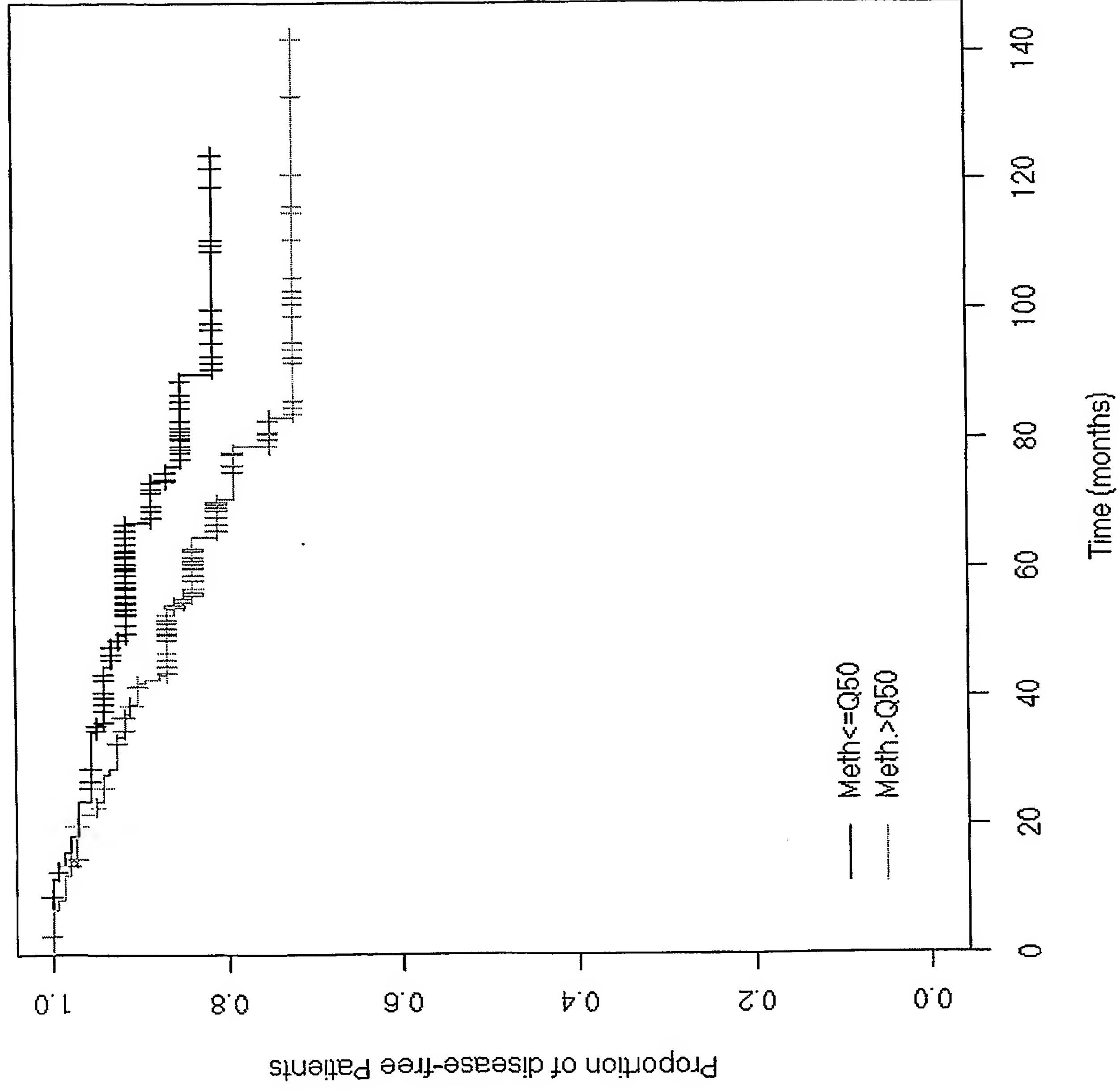


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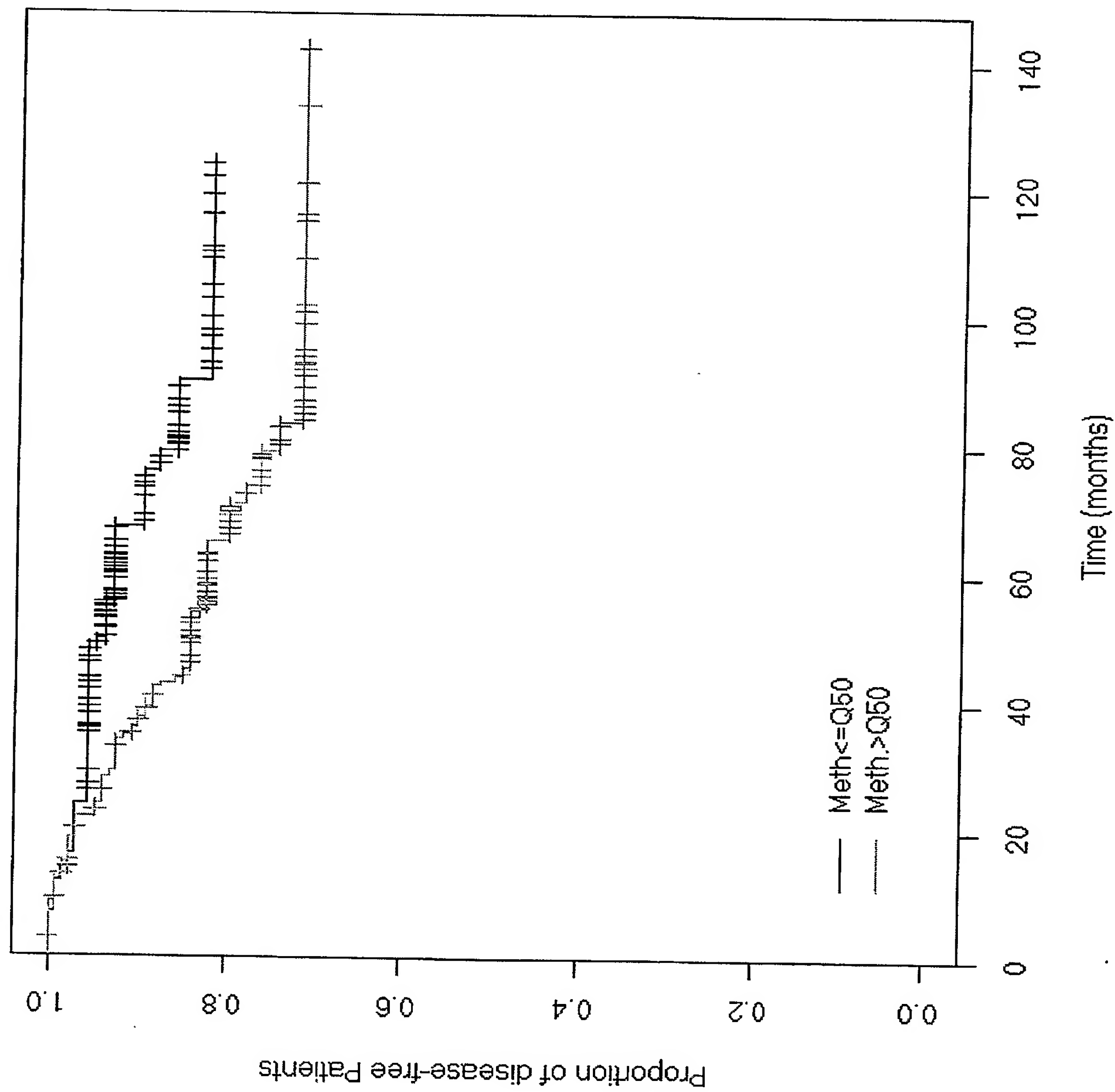


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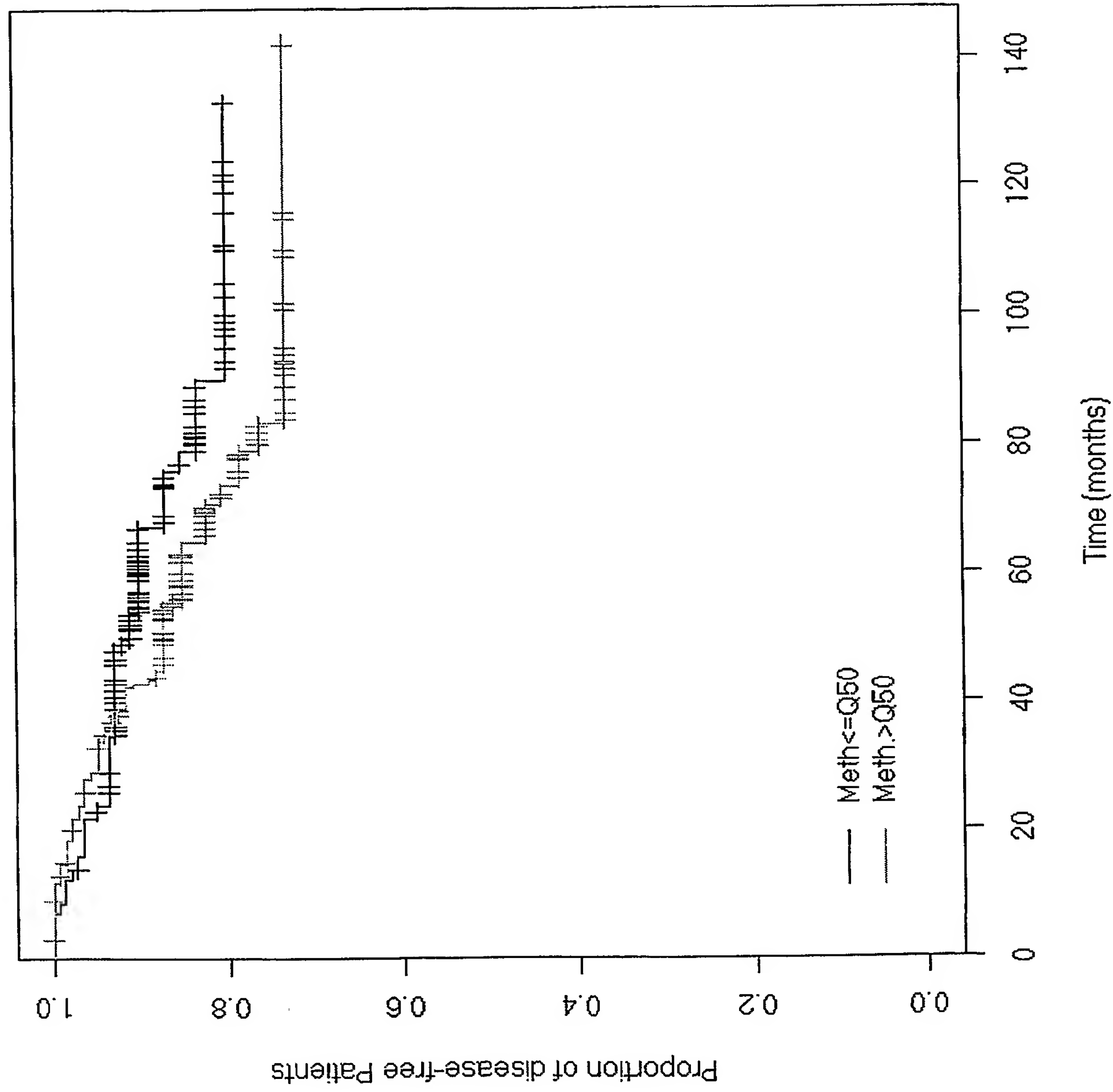


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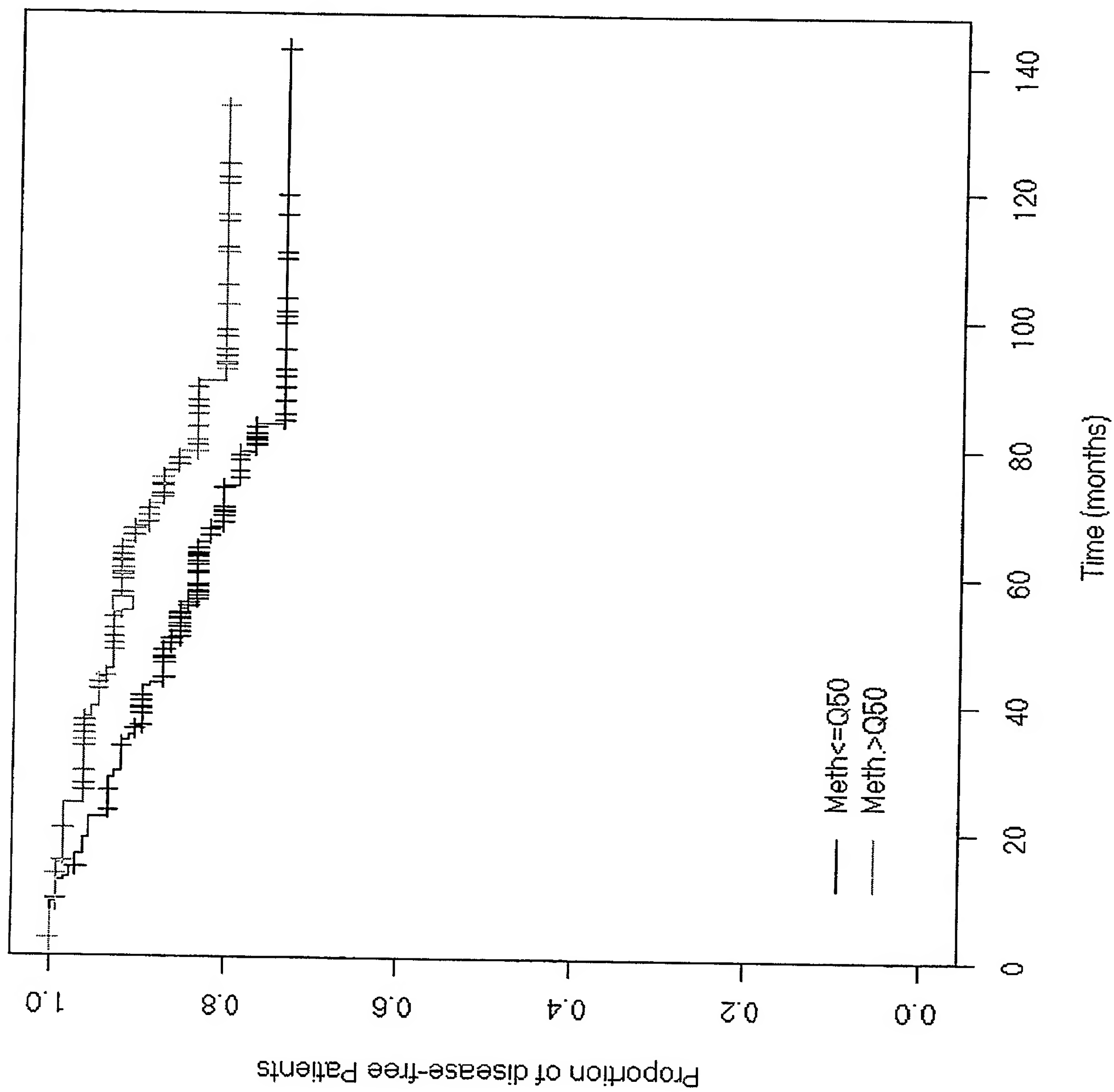


Figure 25 SEQ ID NO: 1082

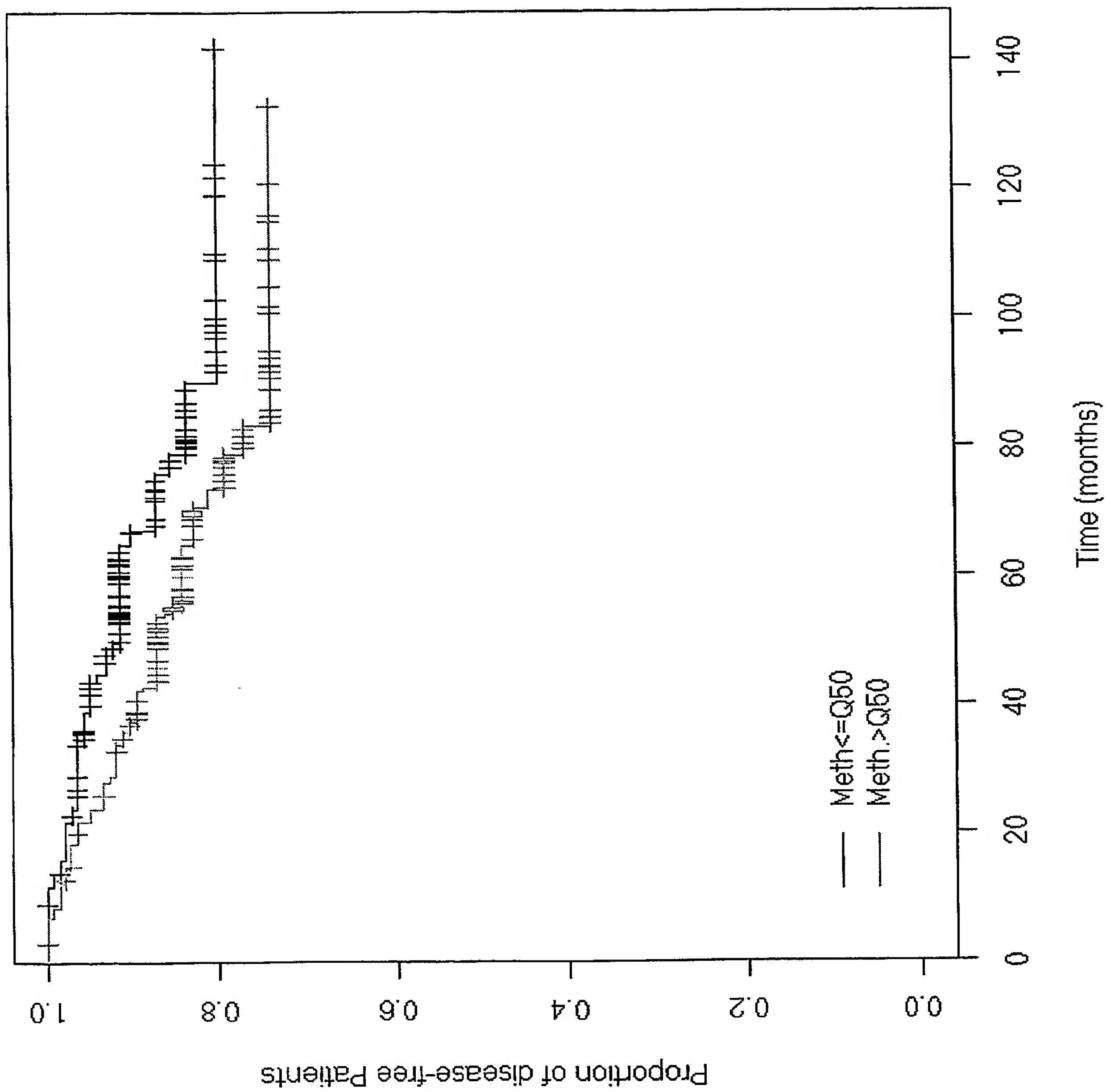


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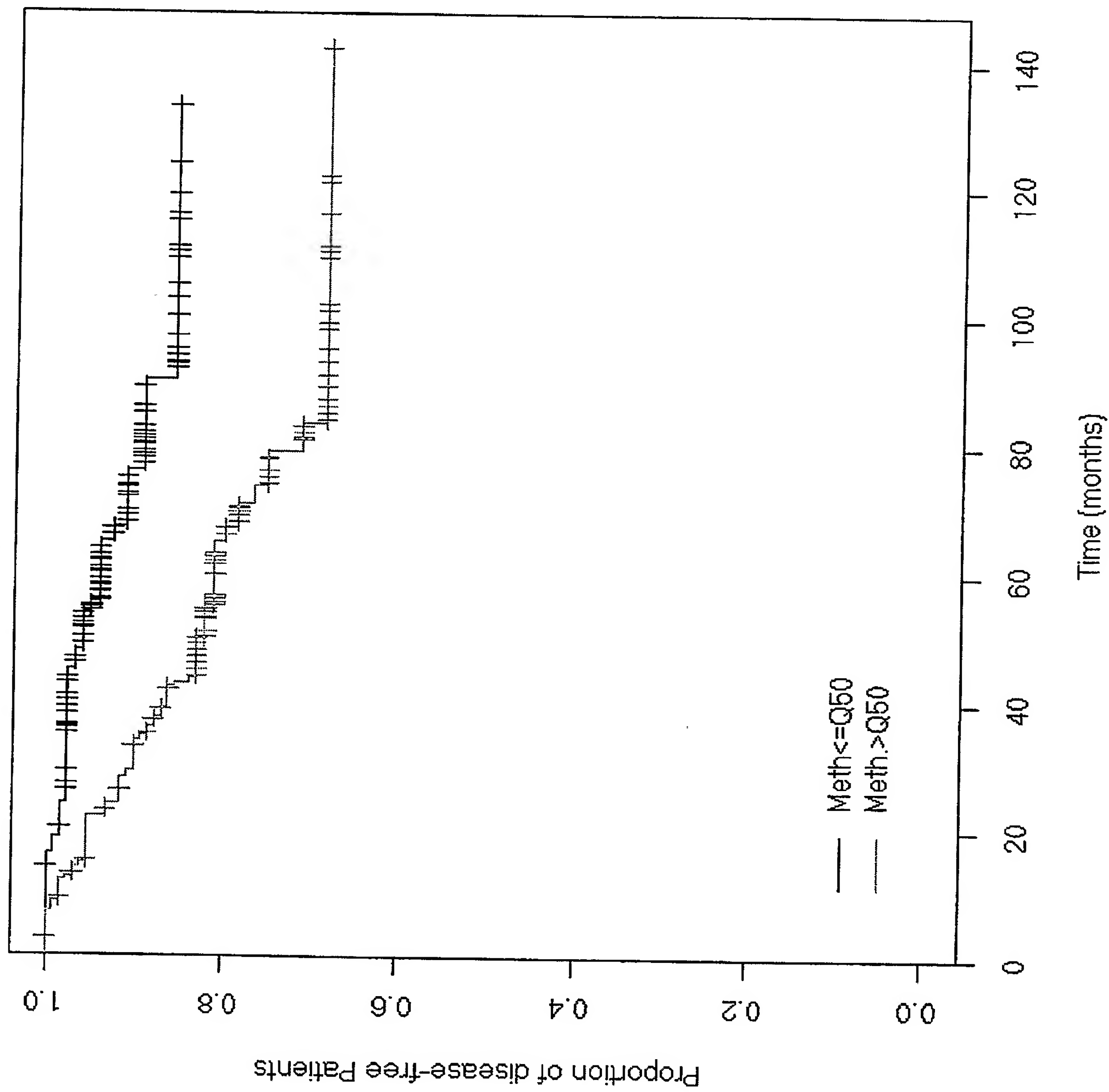


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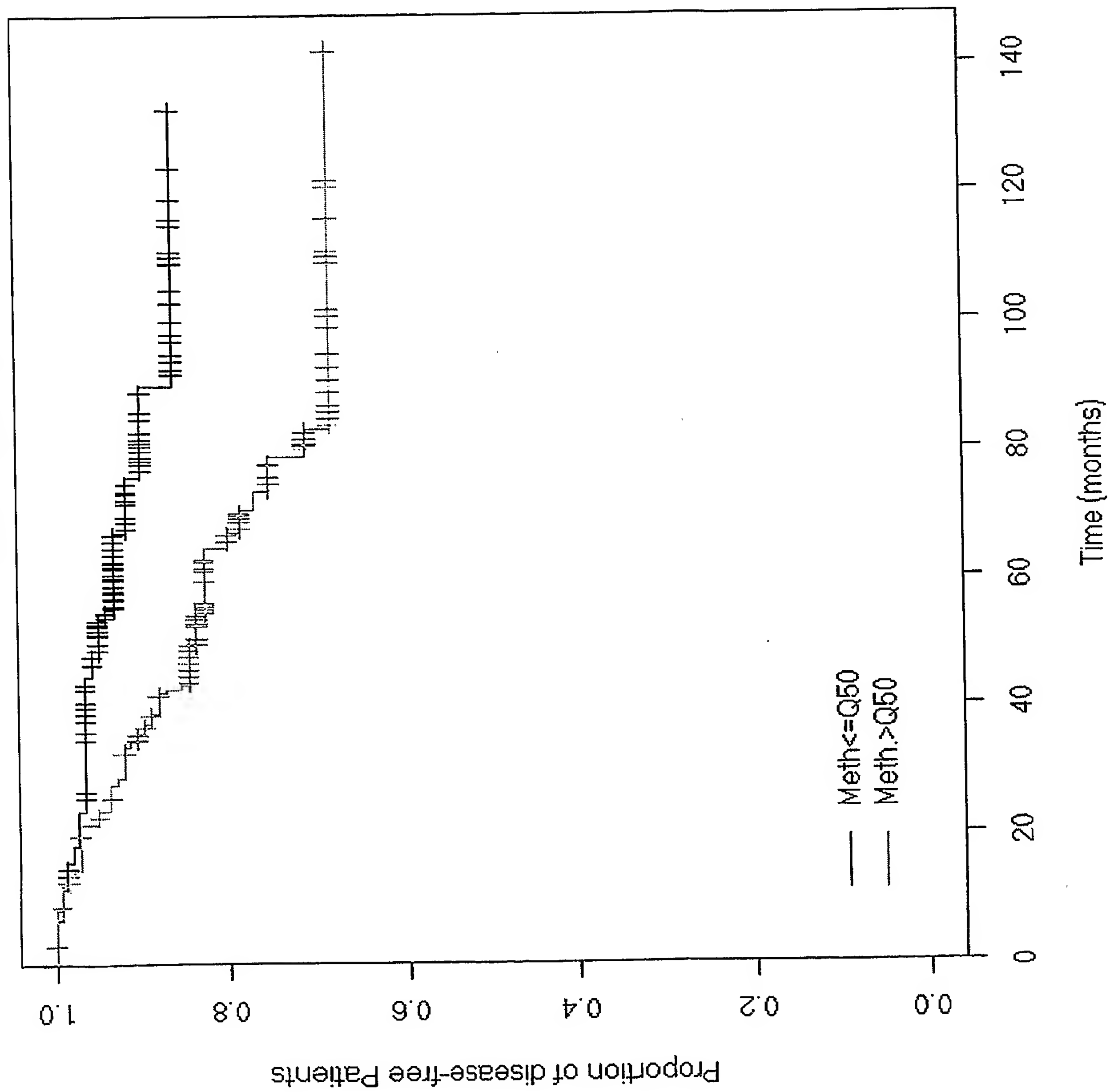


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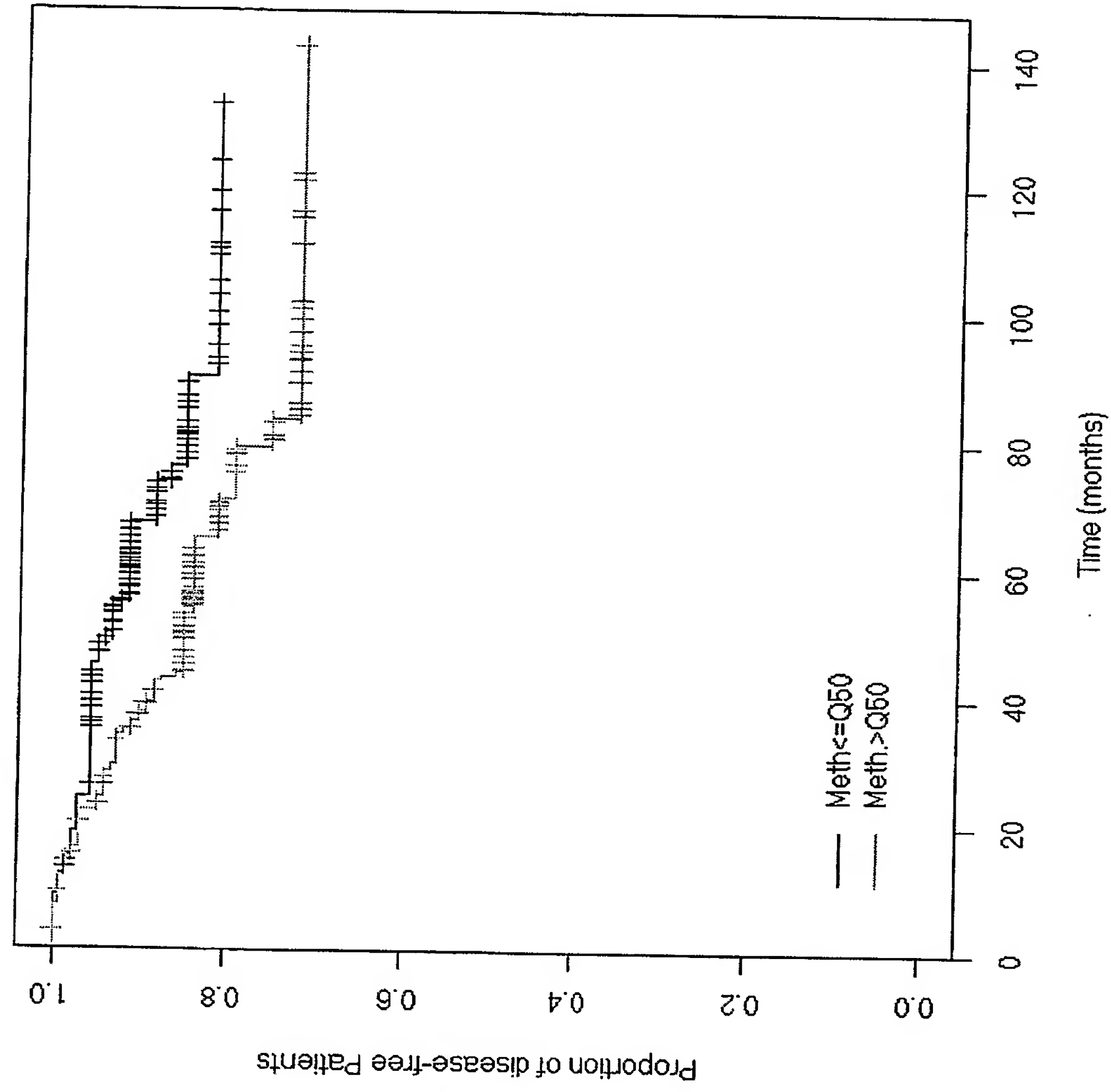


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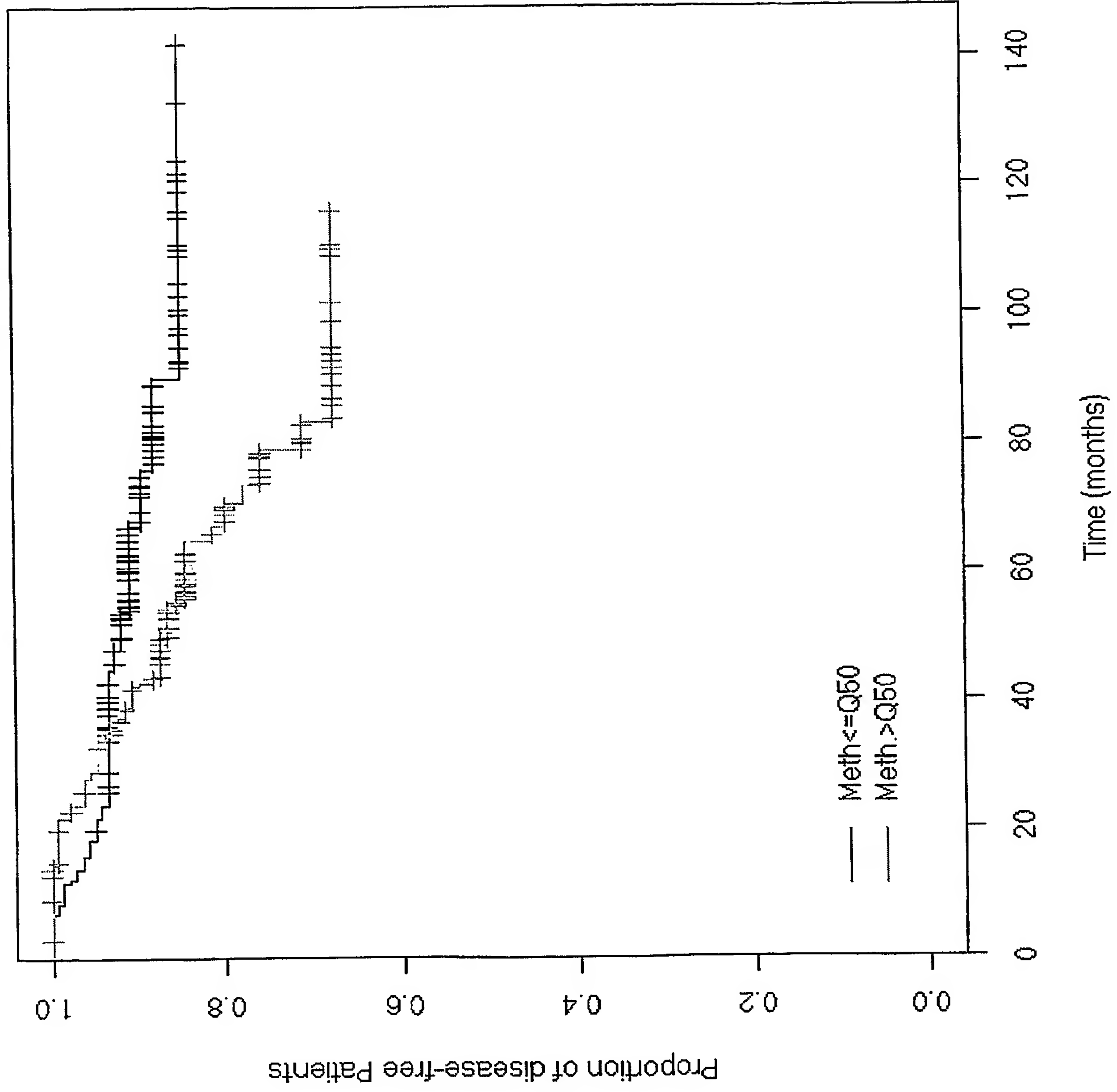


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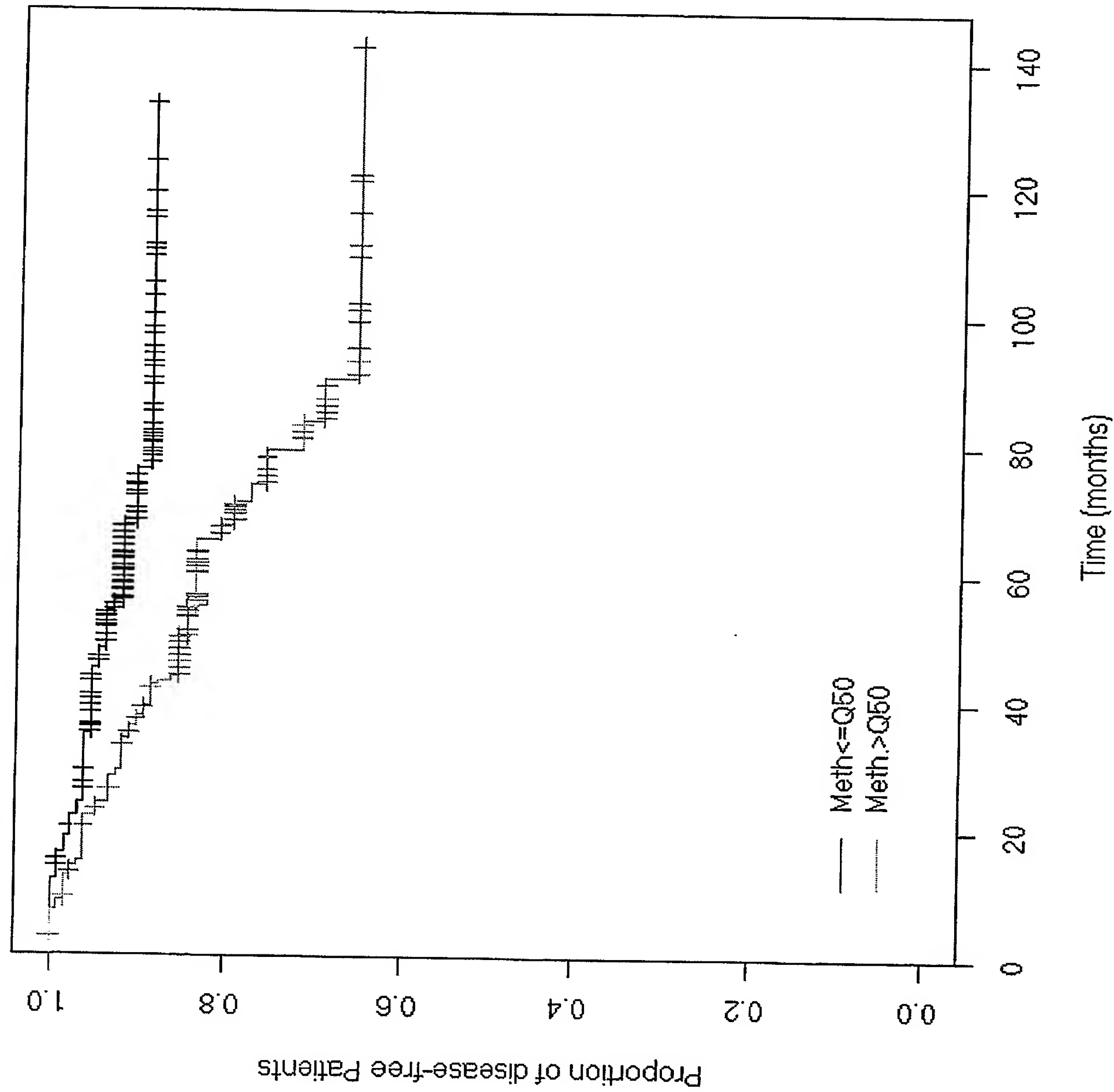


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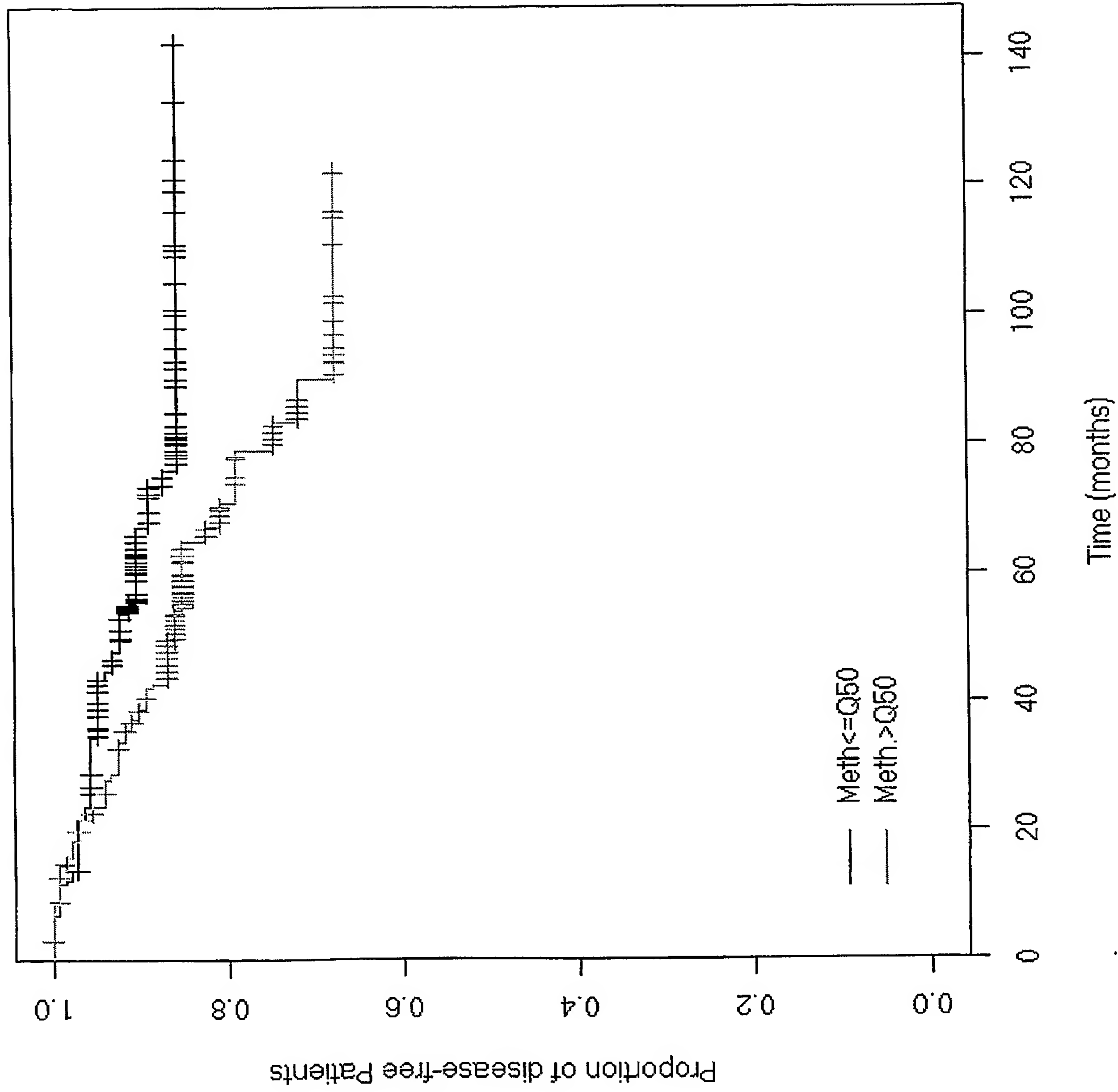


Figure 32 SEQ ID NO: 975

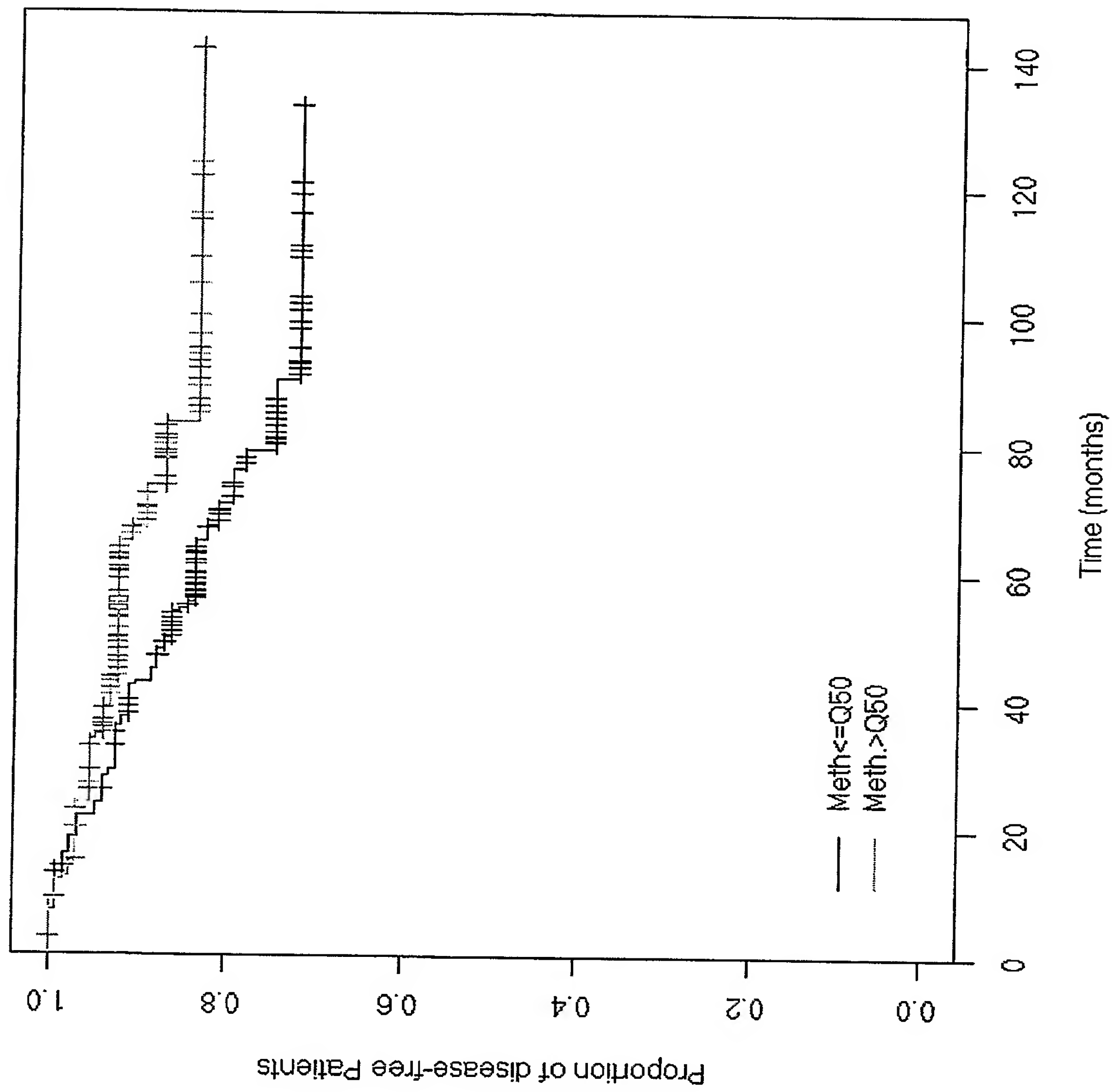


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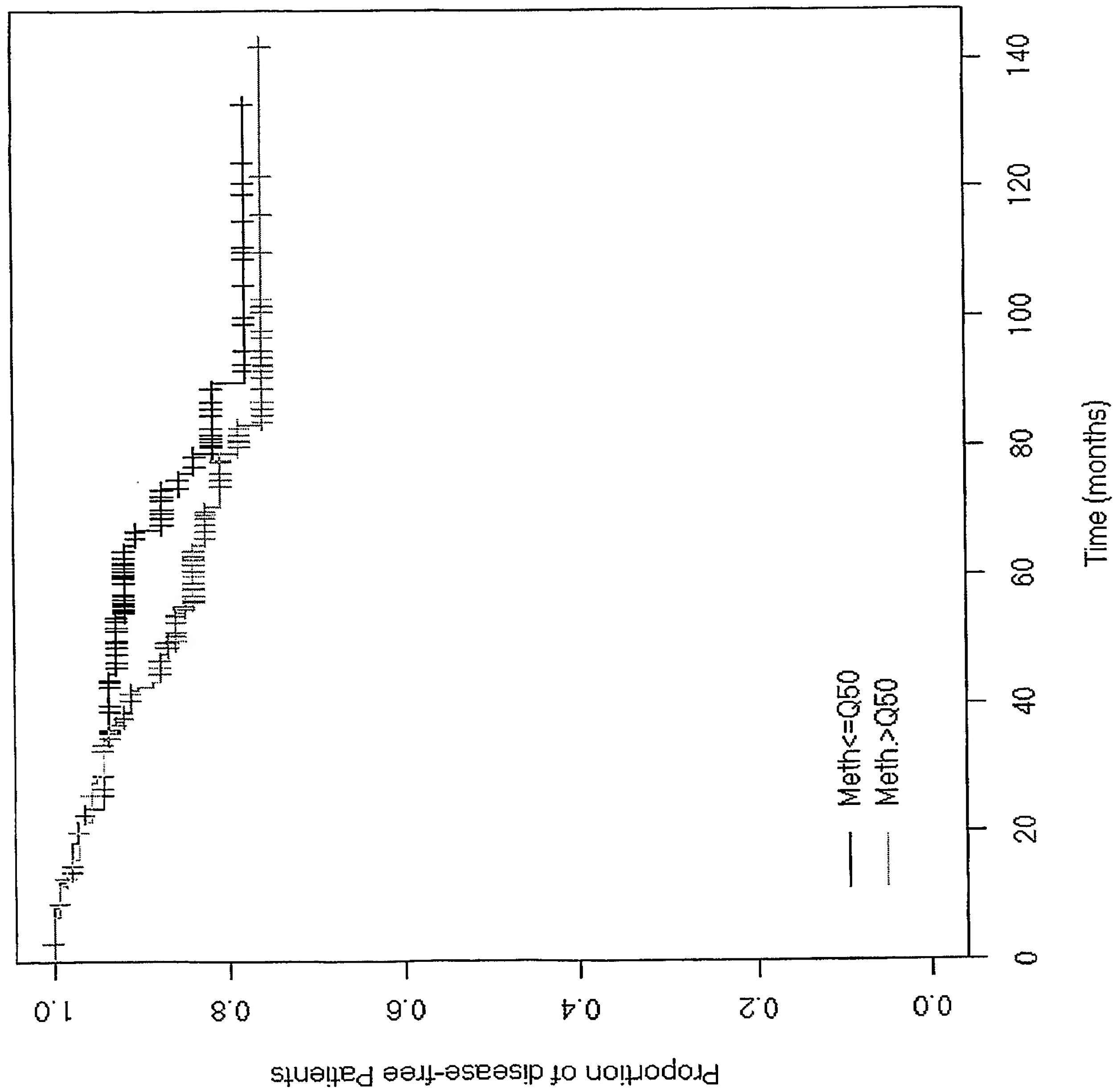


Figure 34 SEQ ID NO: 866

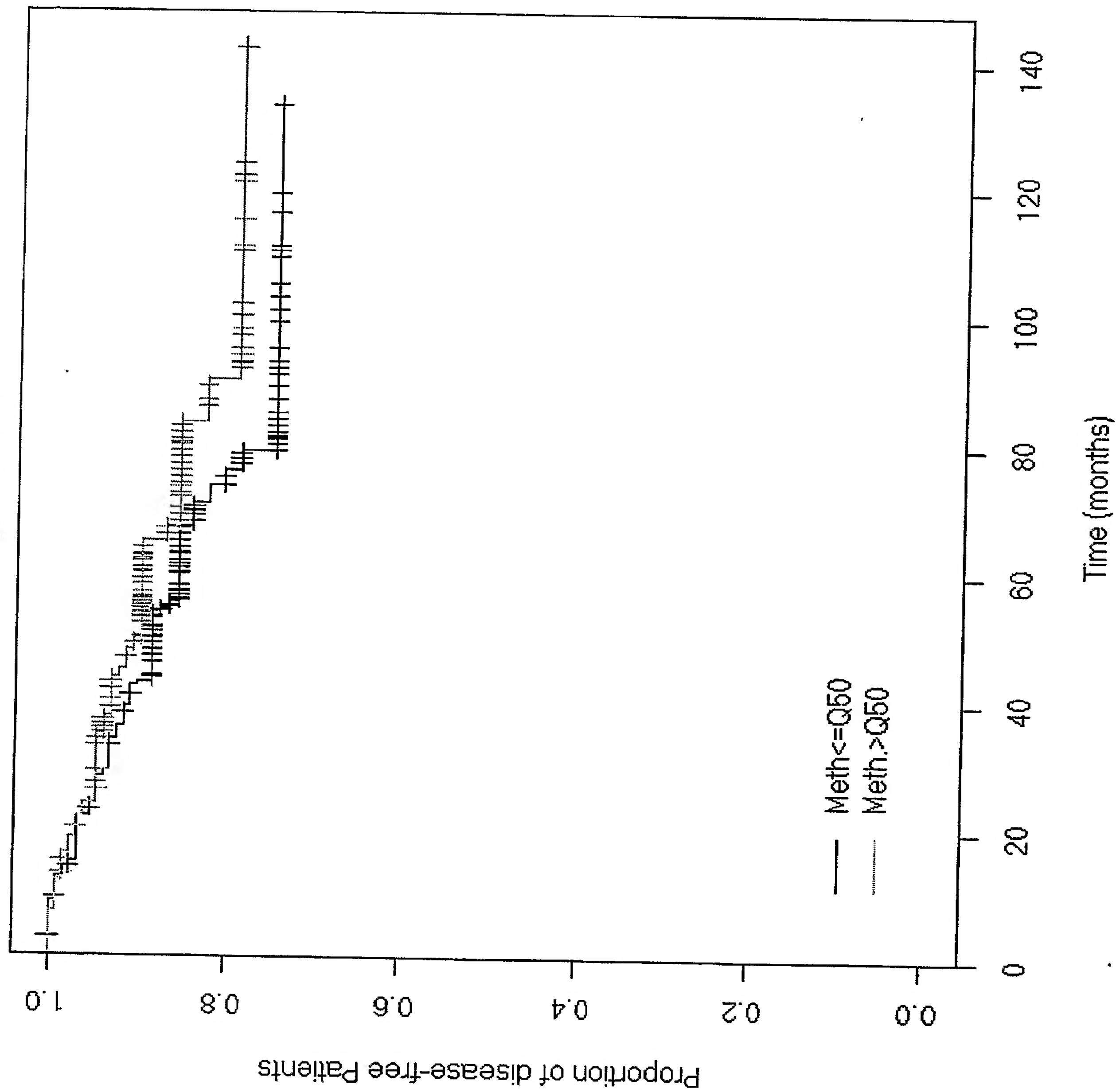


Figure 35

Marker ABCA8 (N= 278)

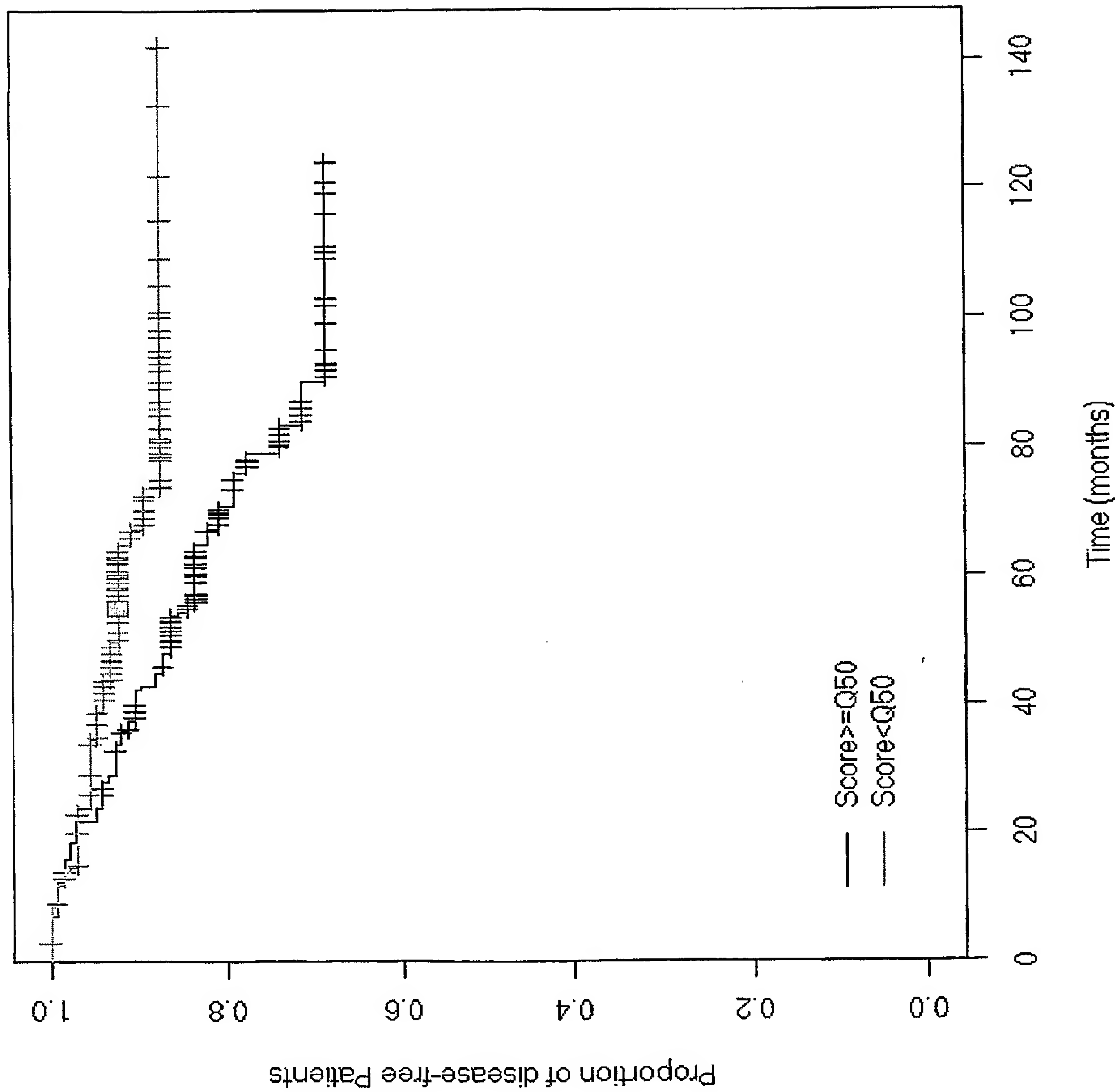


Figure 36 Marker BCL6 (N= 278)

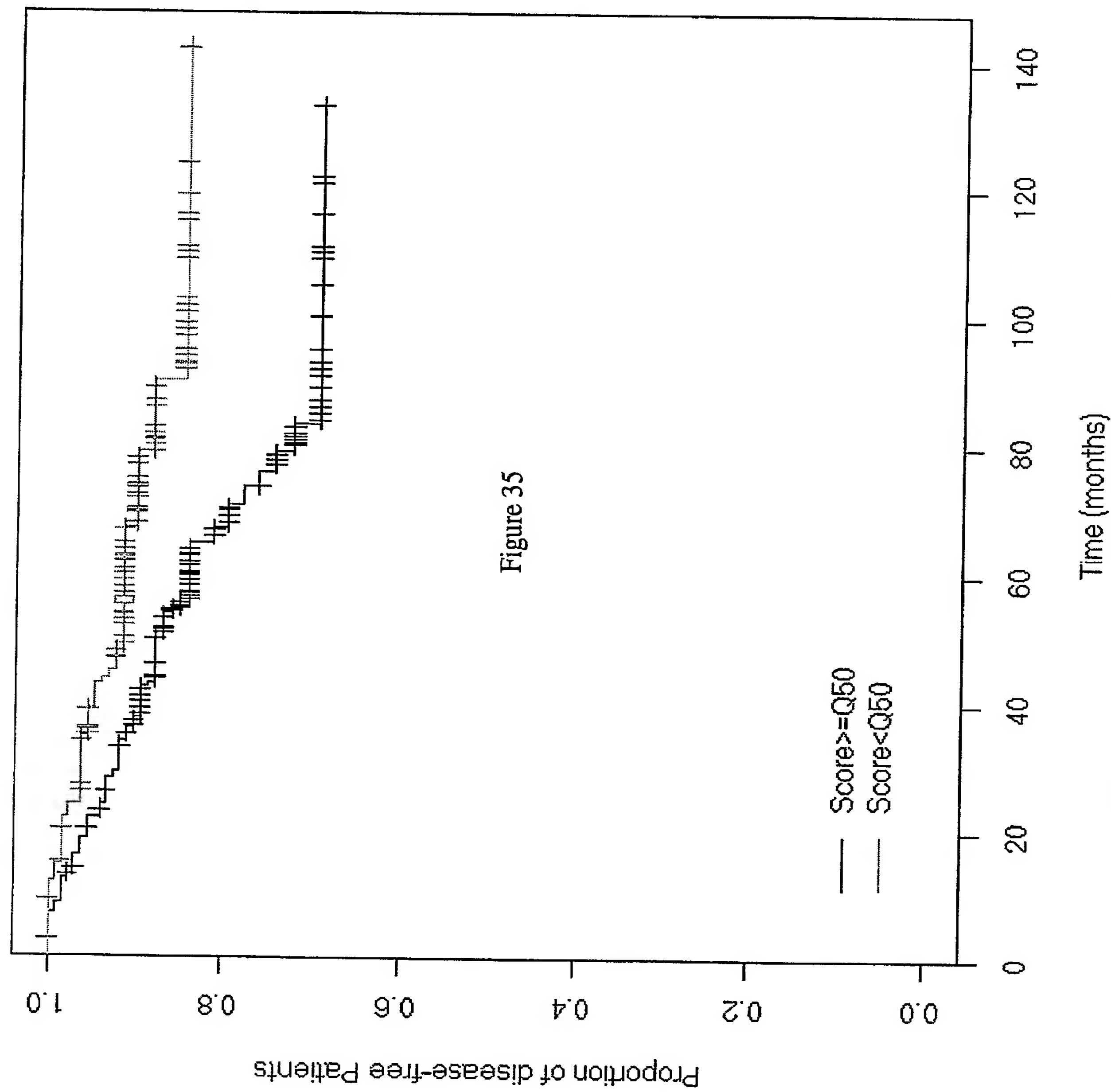


Figure 37

Marker CDK6 (N= 278)

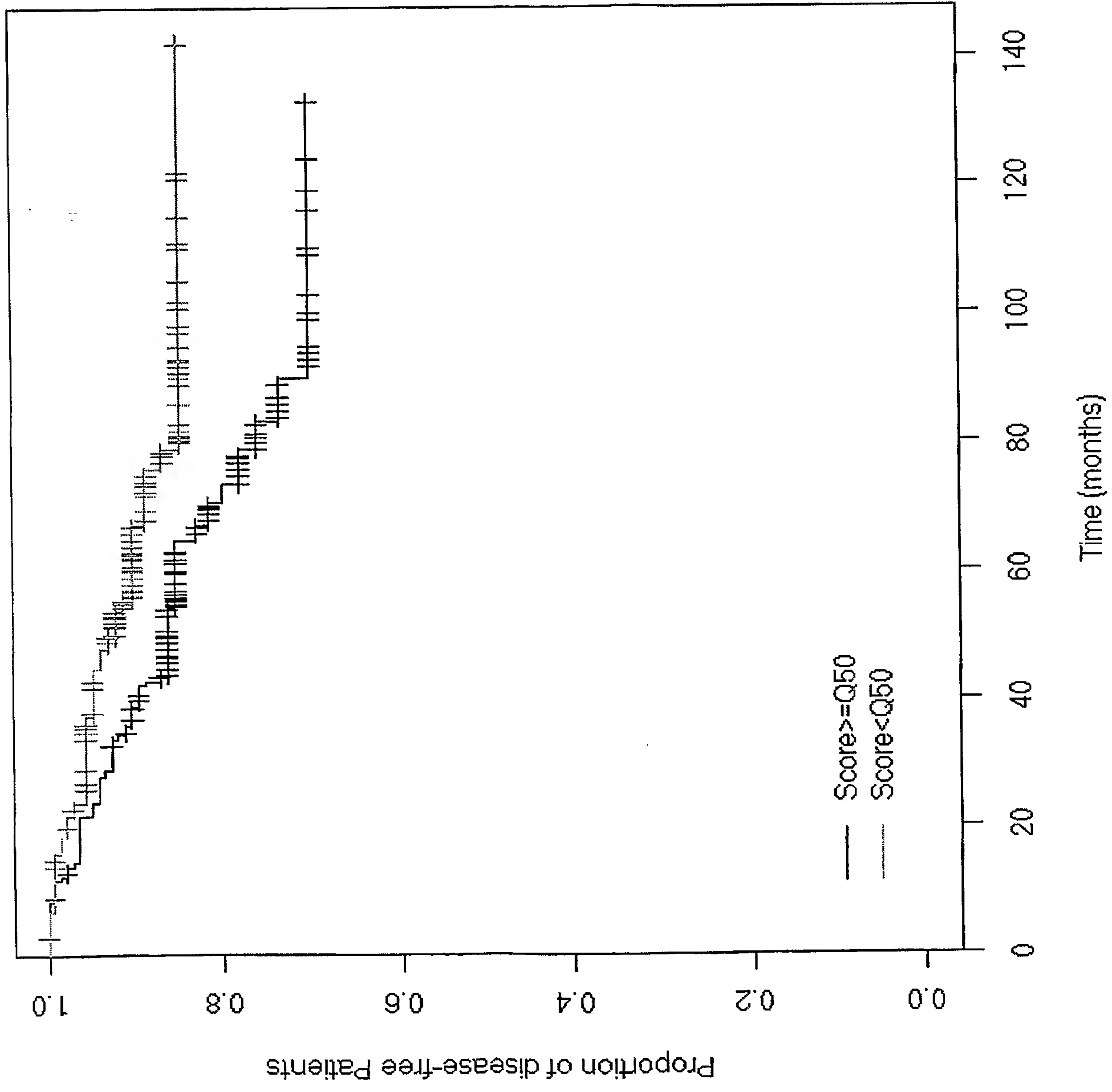
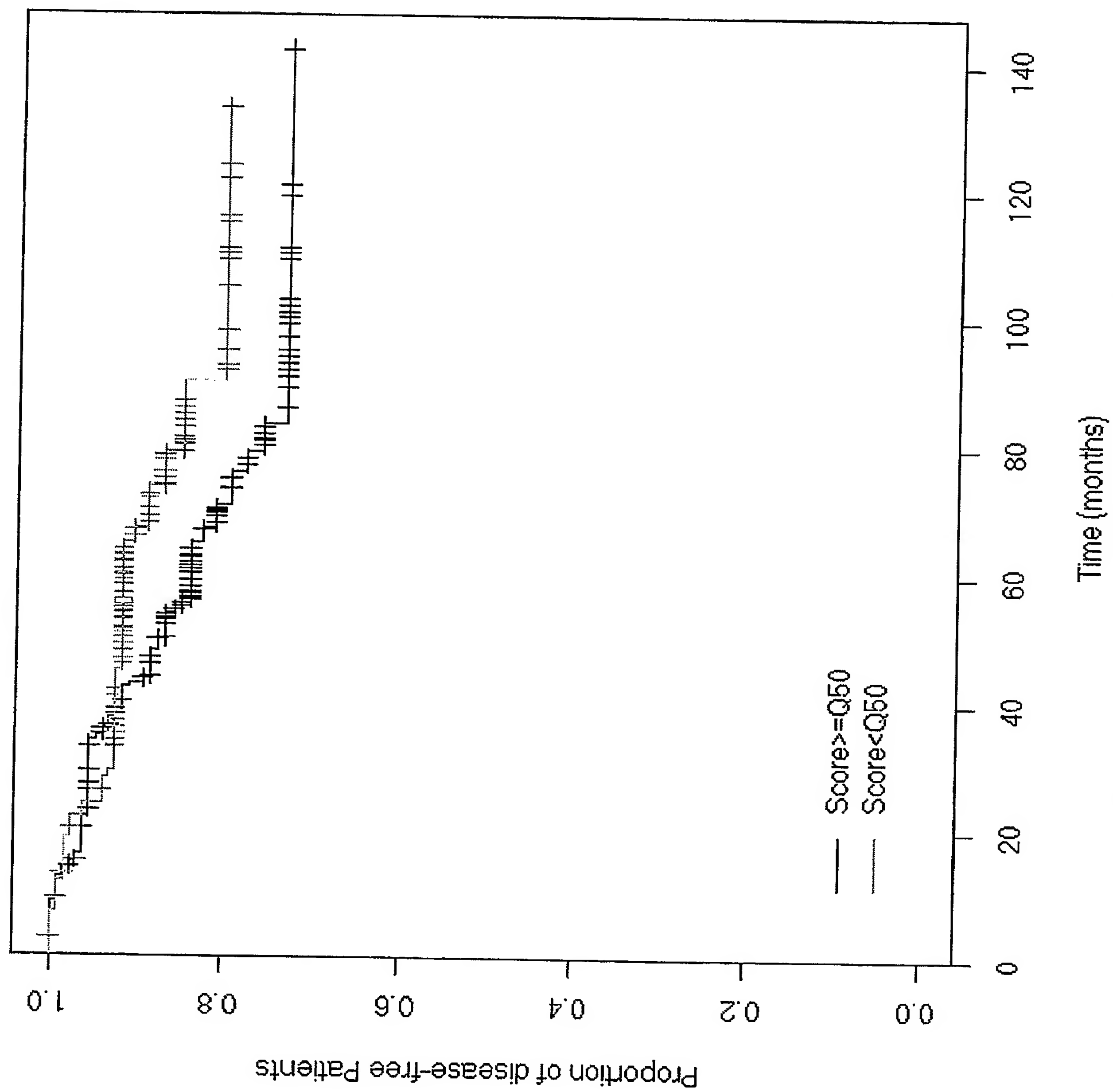


Figure 38 Marker CGB1 (N= 278)



Marker ERBB2 (N= 278)

Figure 39

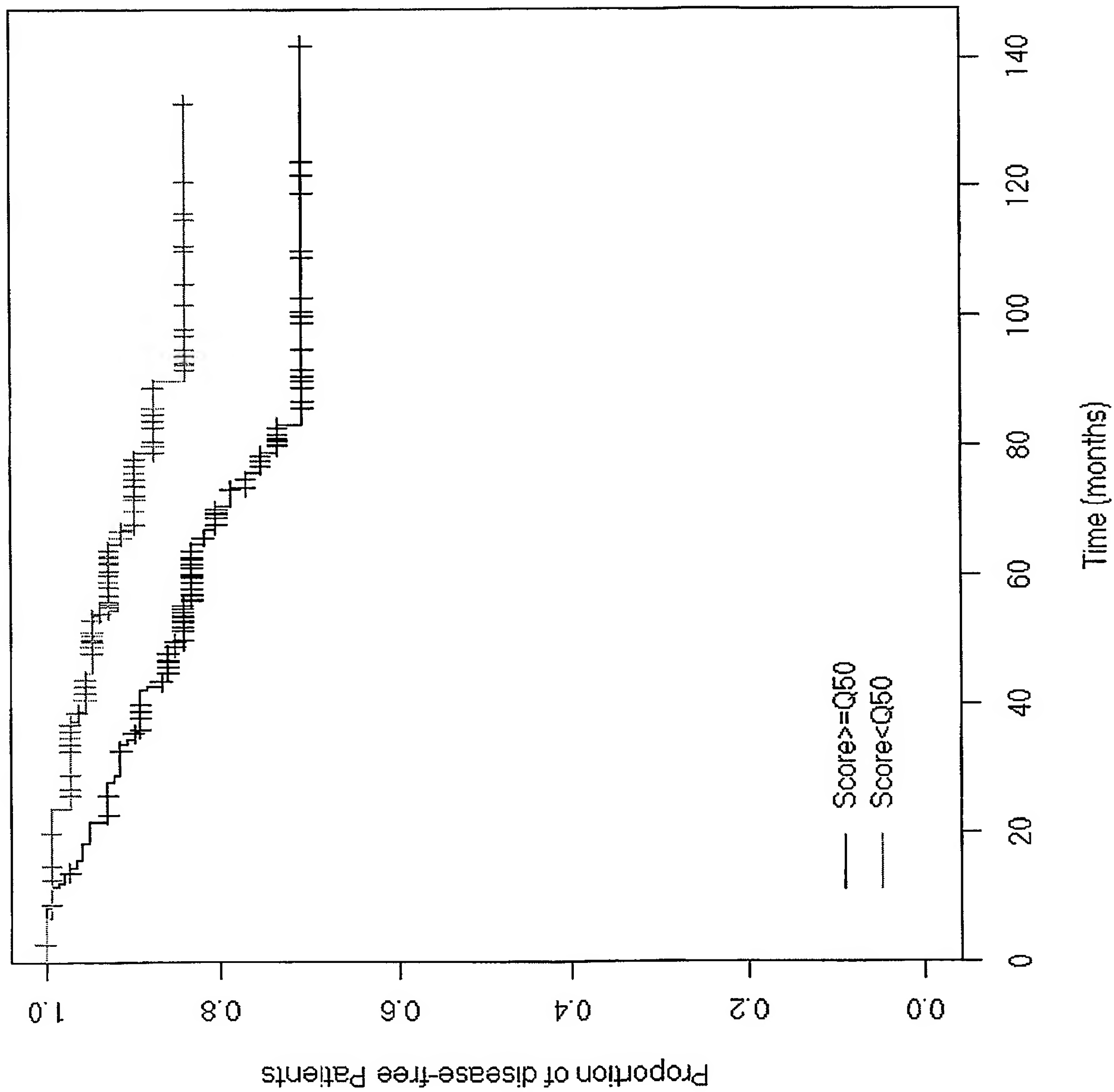


Figure 40

Marker ONECUT2 (N= 278)

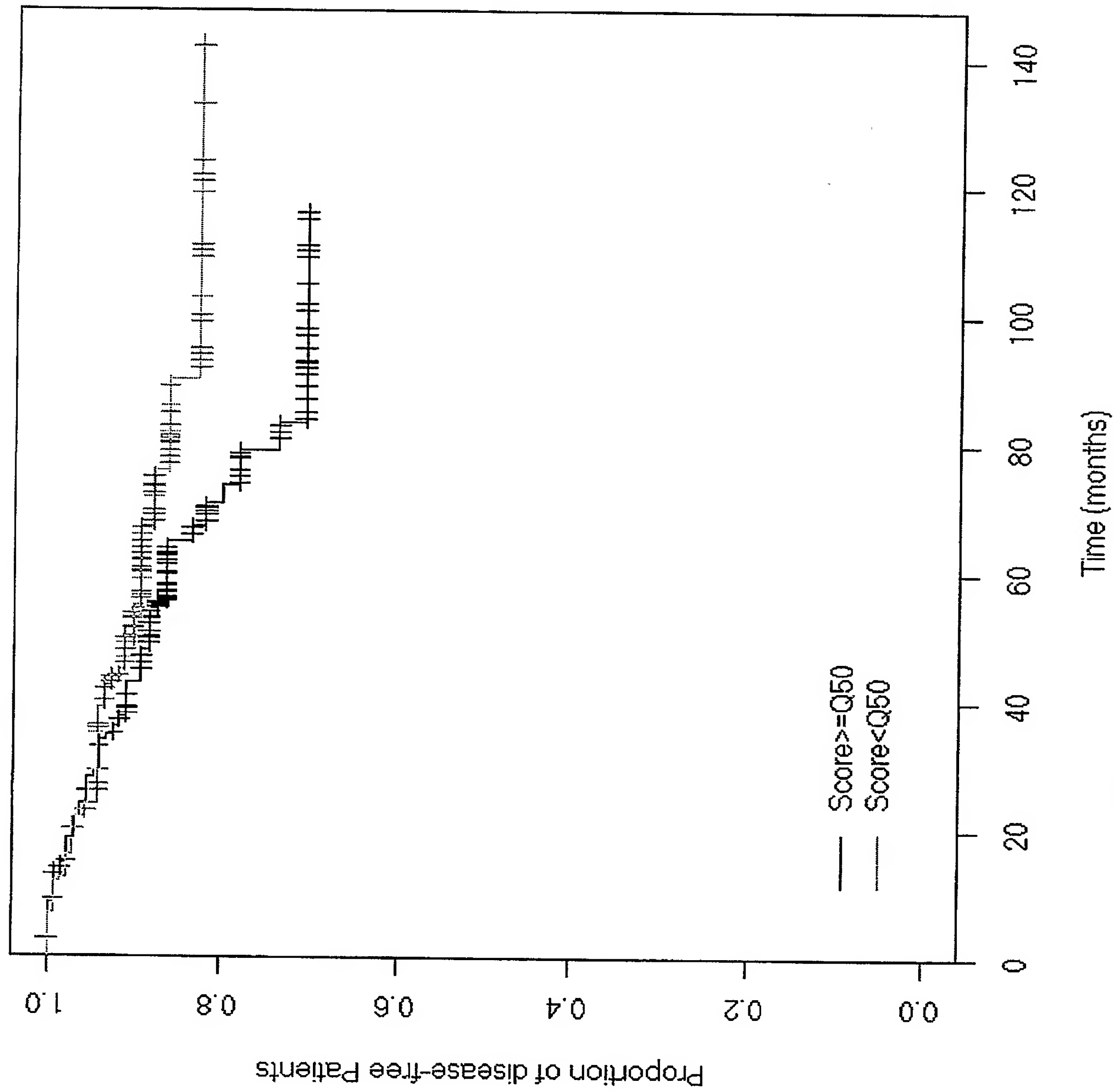


Figure 41

Marker PITX2 (N= 278)

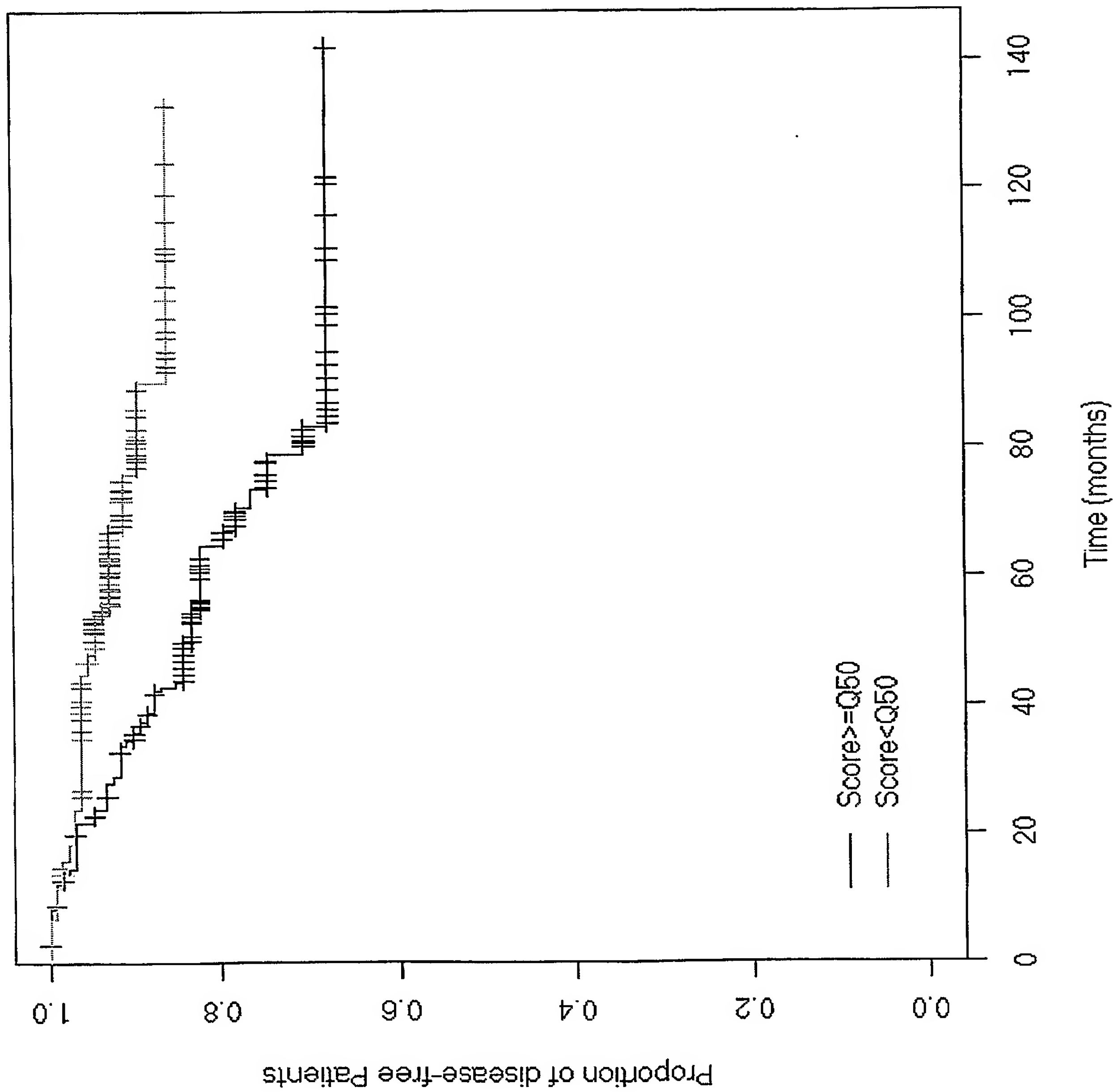


Figure 42 **Marker PLAU (N= 278)**

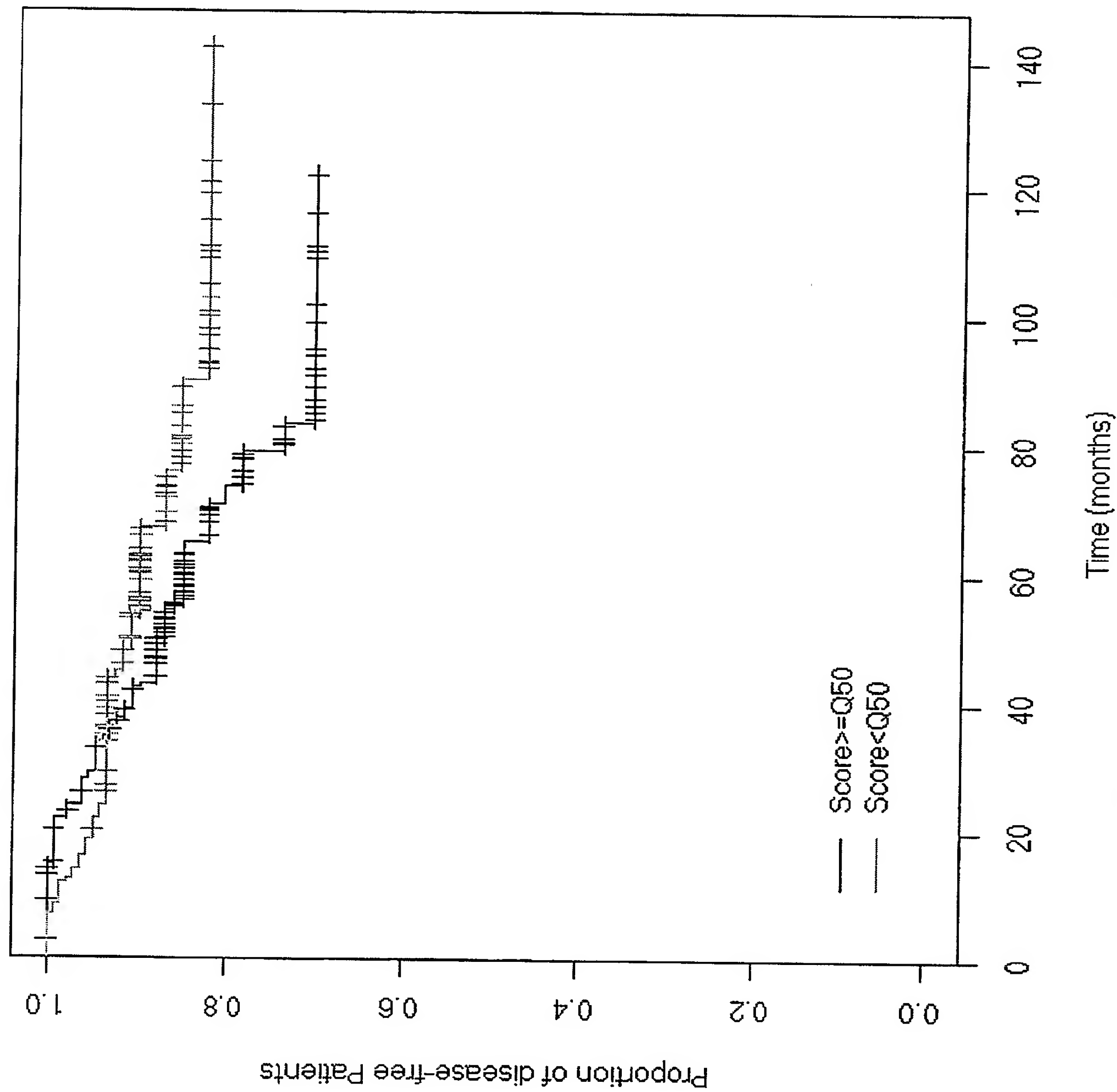


Figure 43

Marker STMN1 (N= 278)

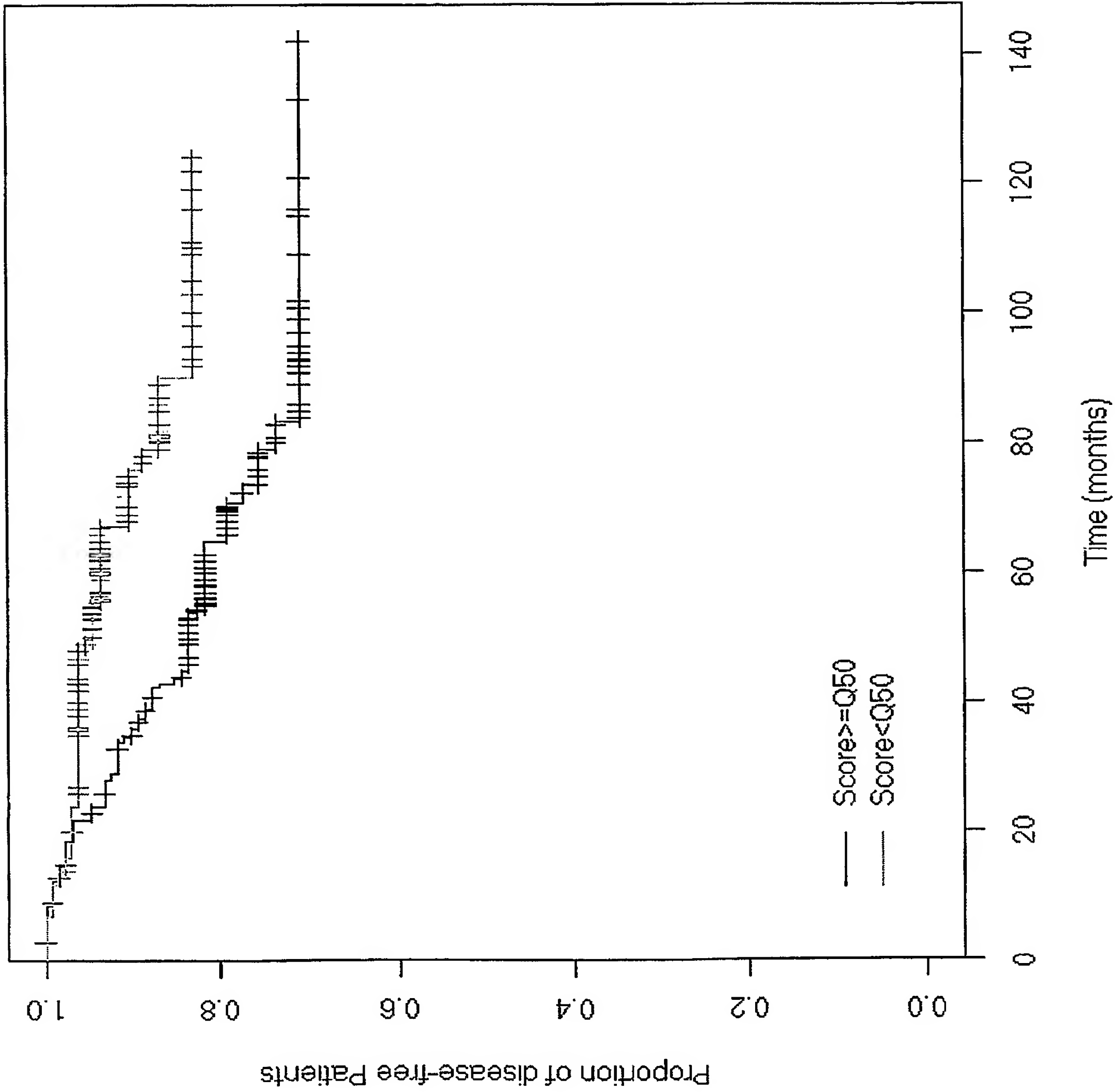


Figure 44
Marker TBC1D3 (N= 278)

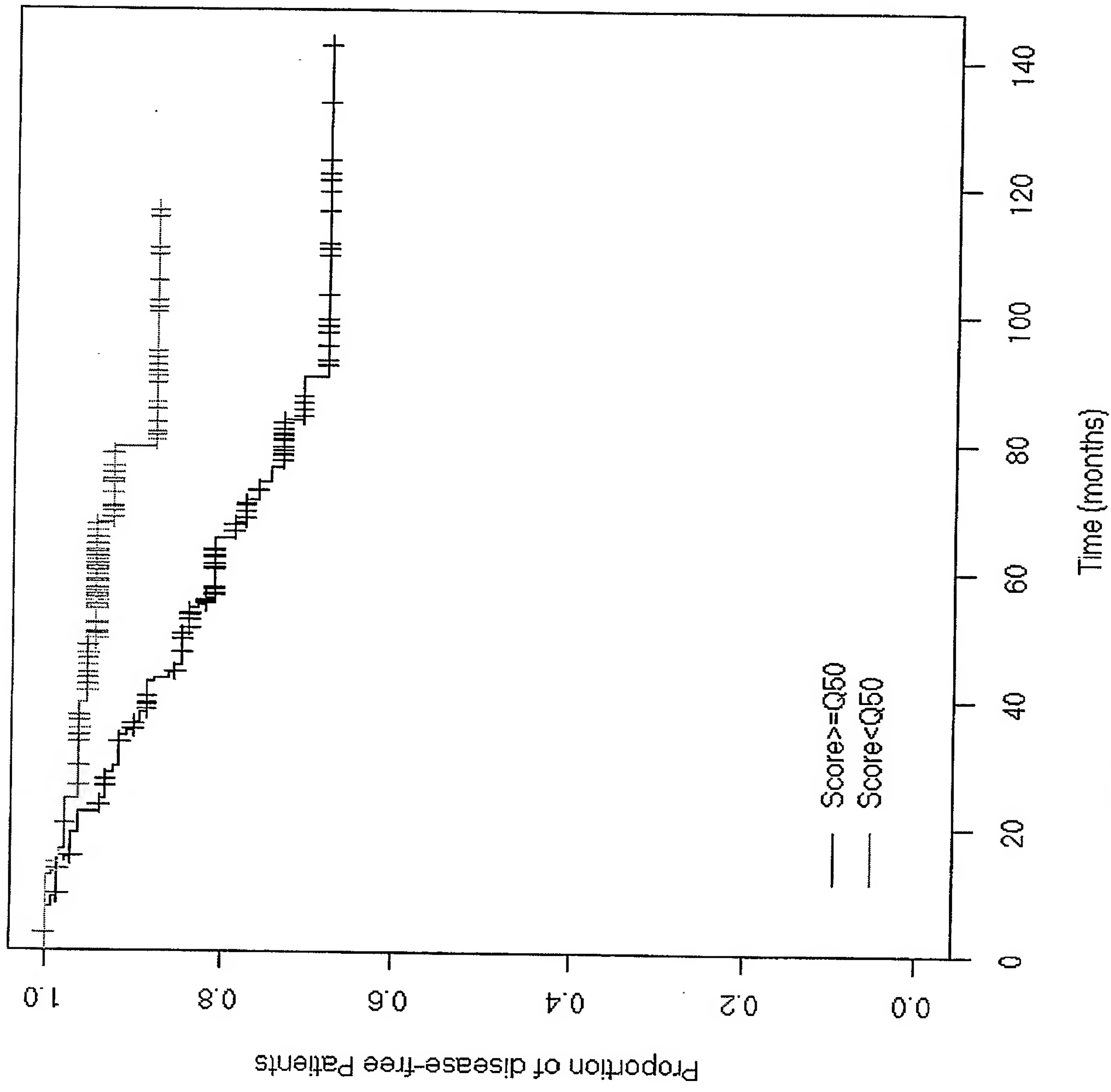


Figure 45

Marker VTN (N= 278)

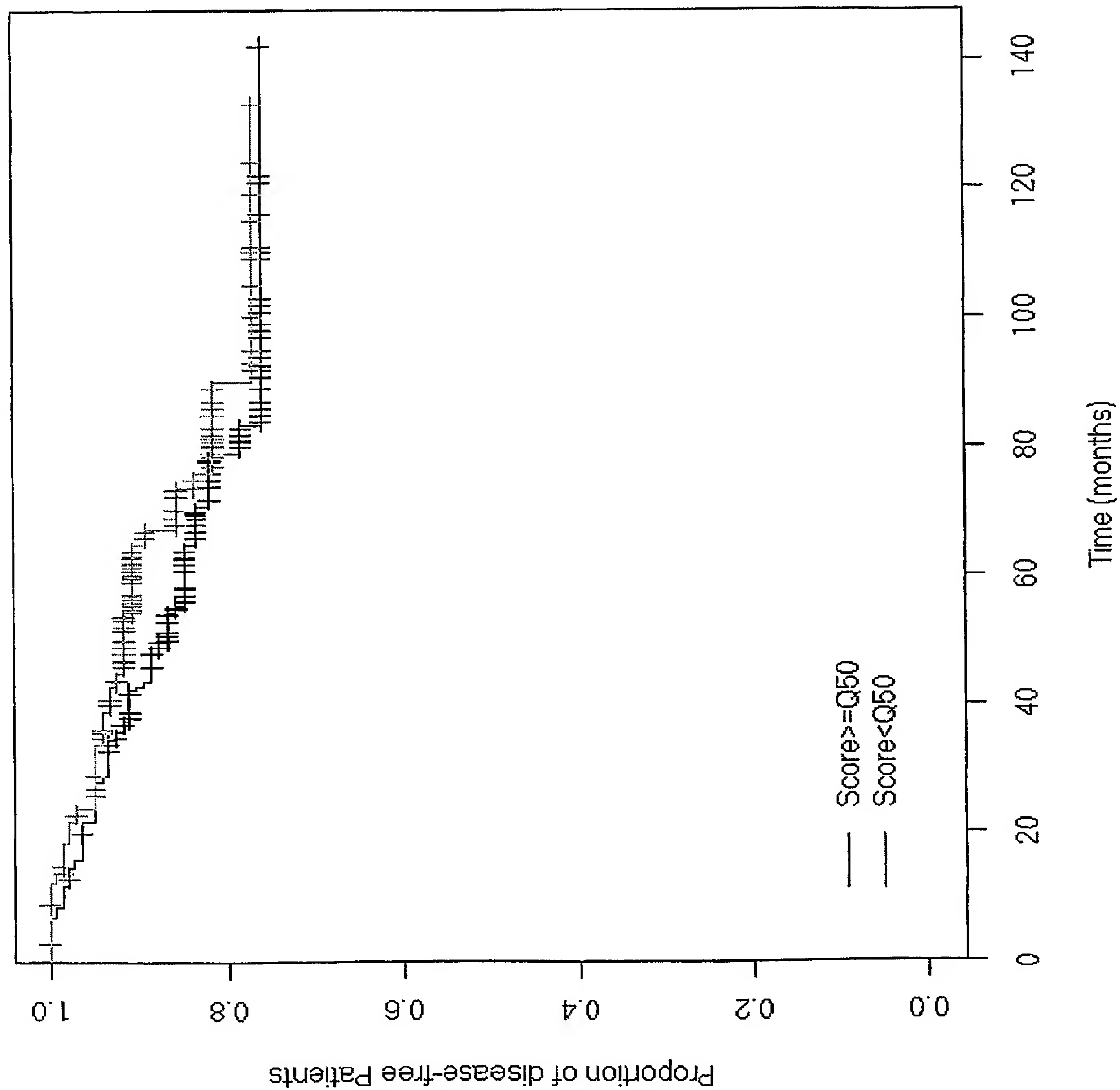
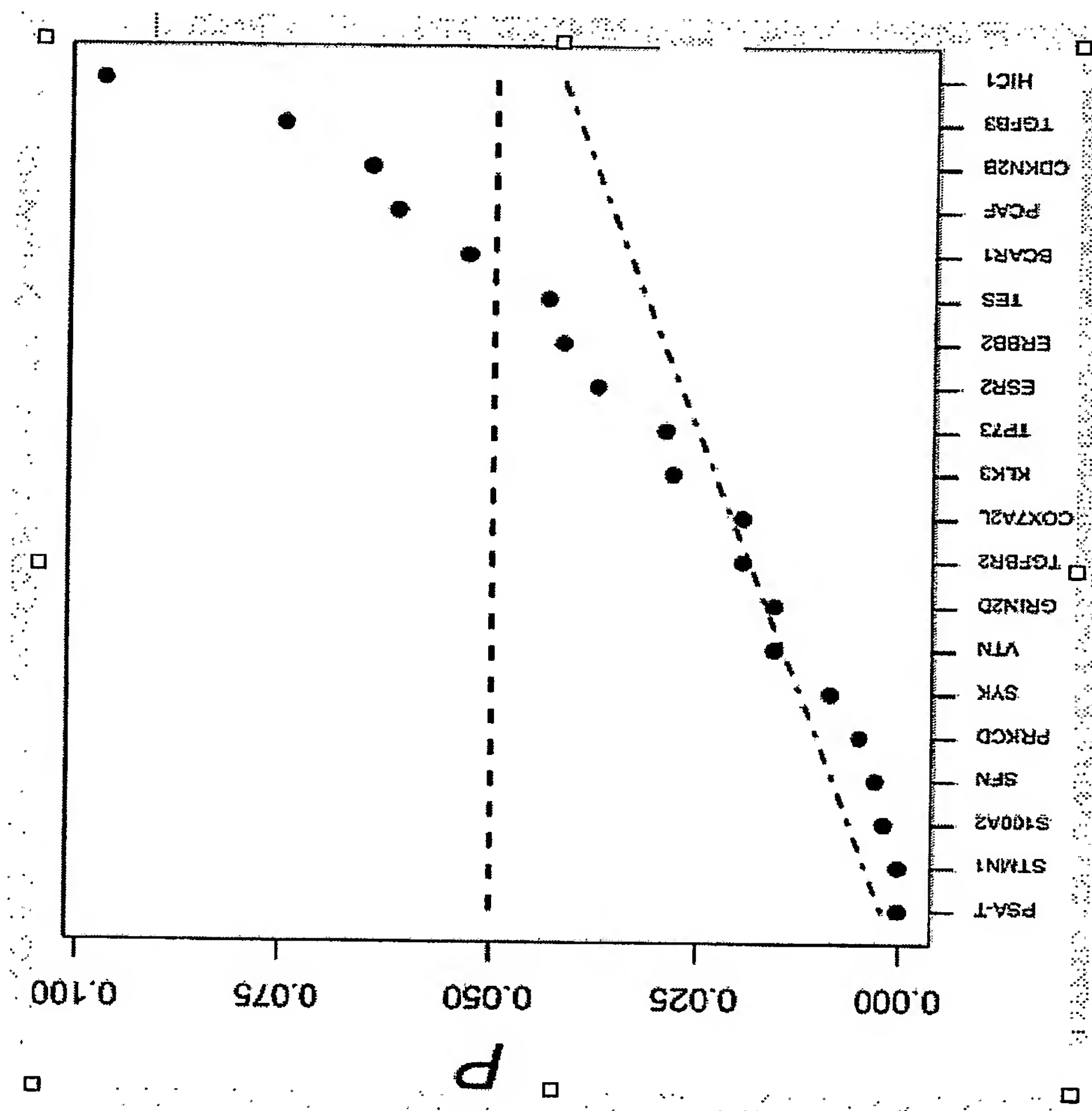


Figure 46



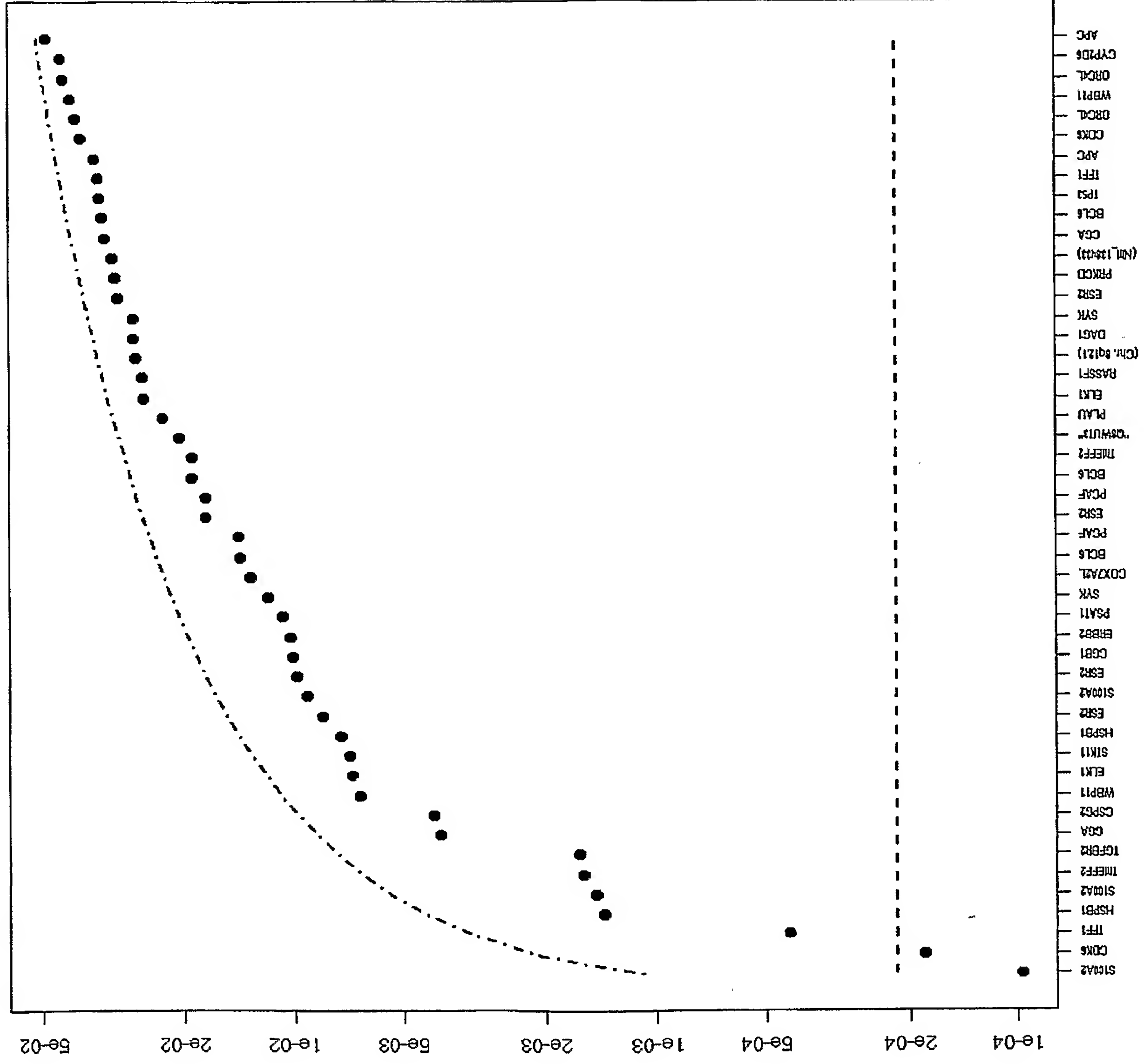
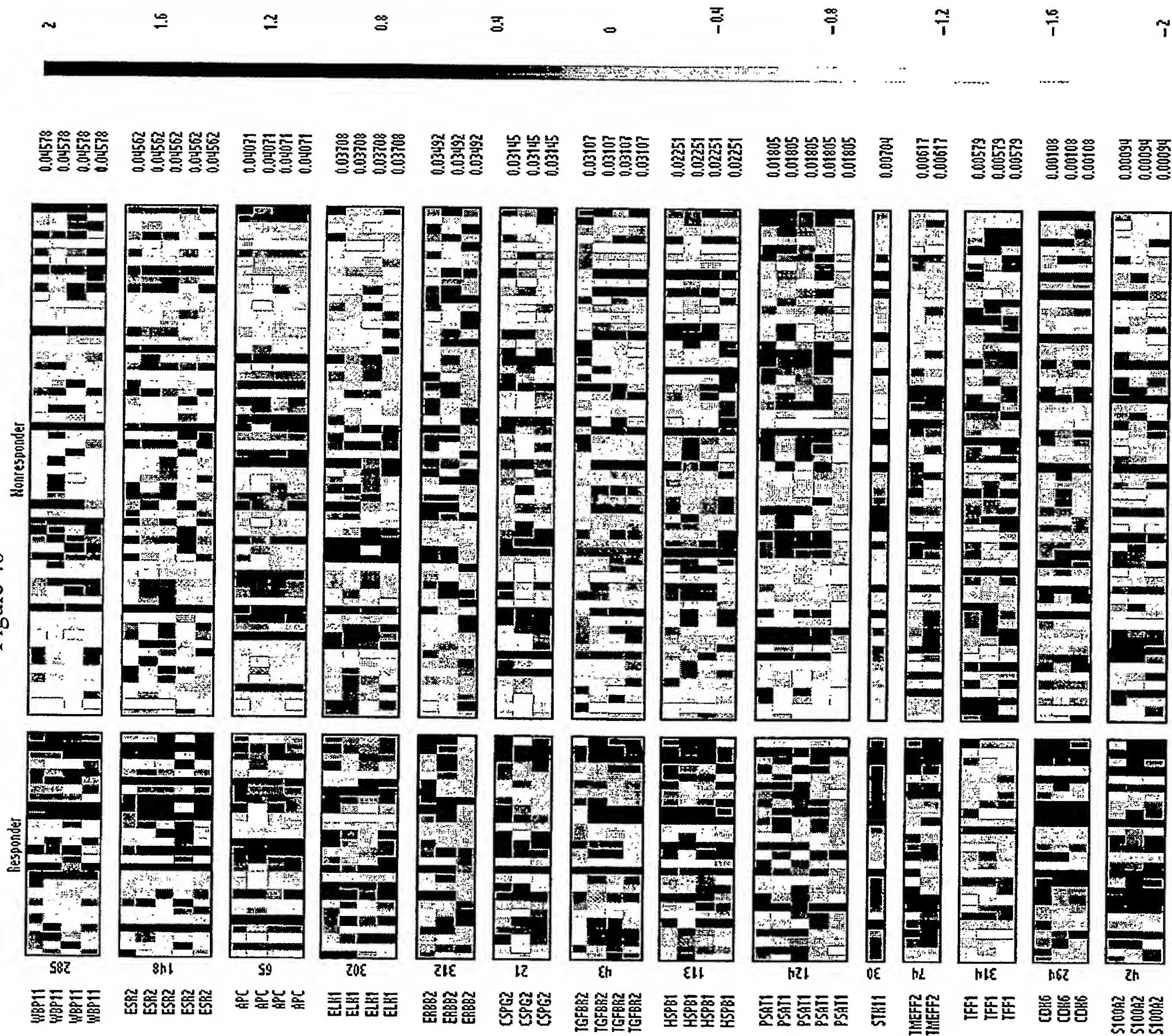


Figure 48



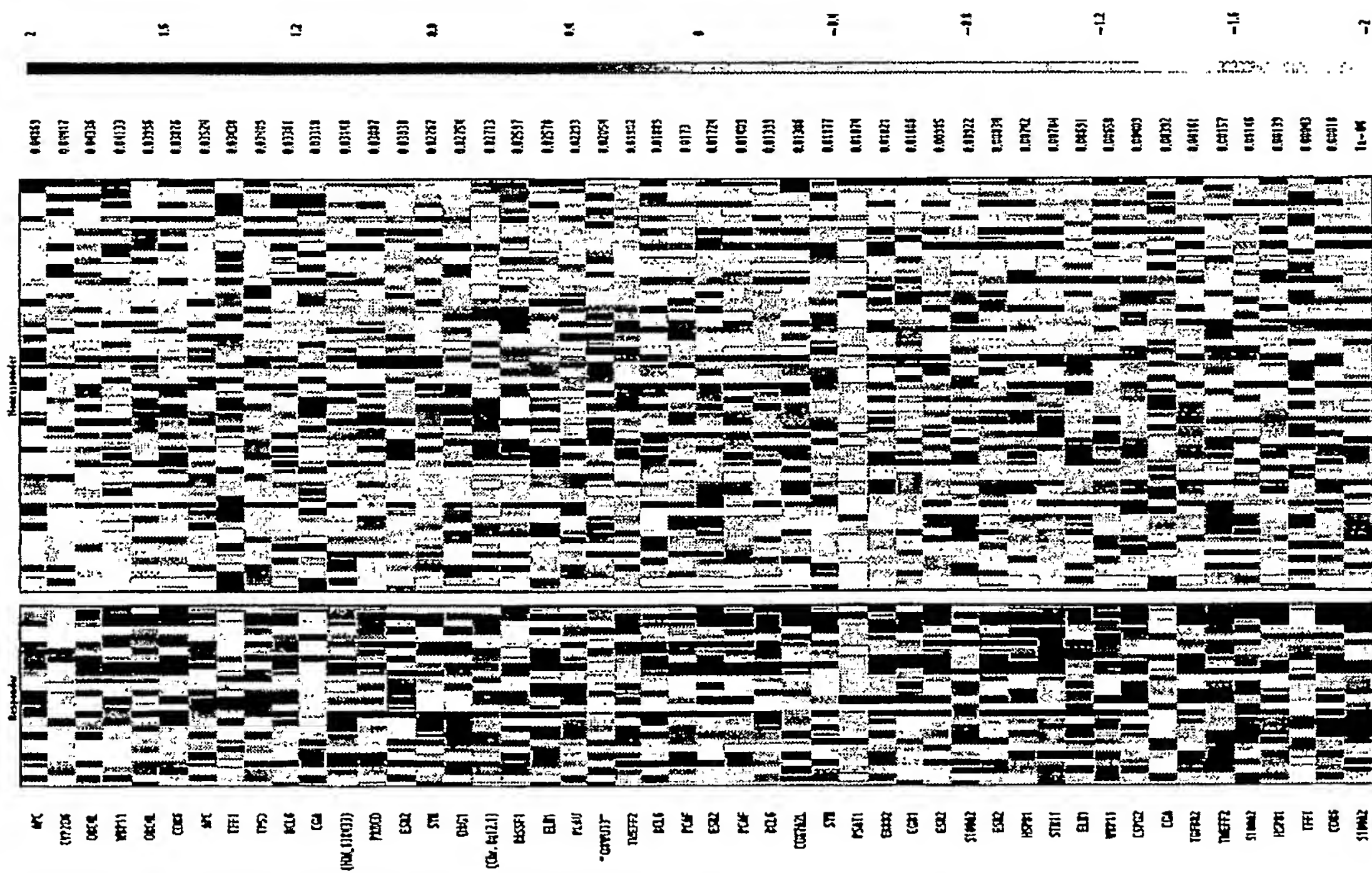


Figure 50

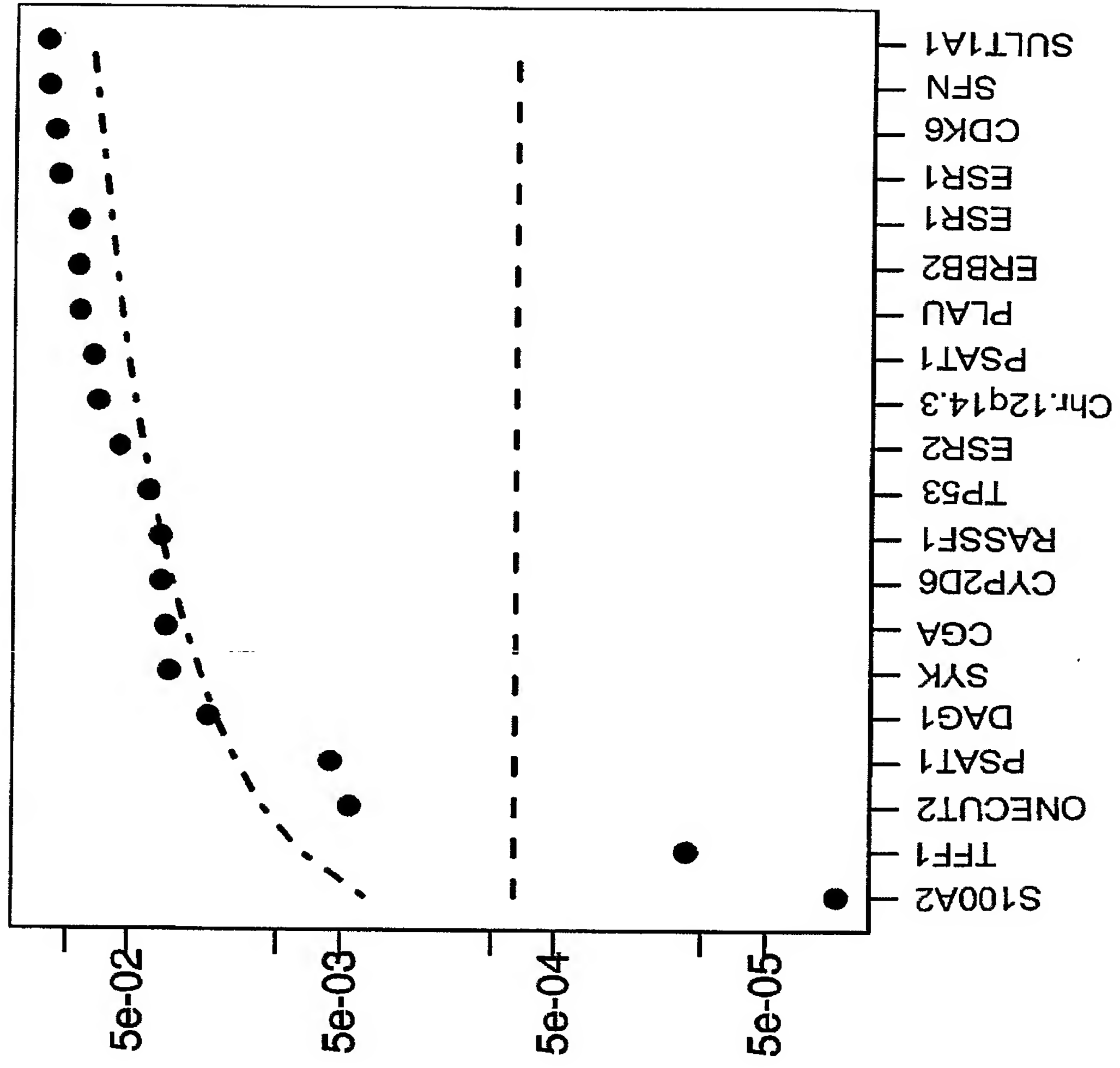


Figure 51

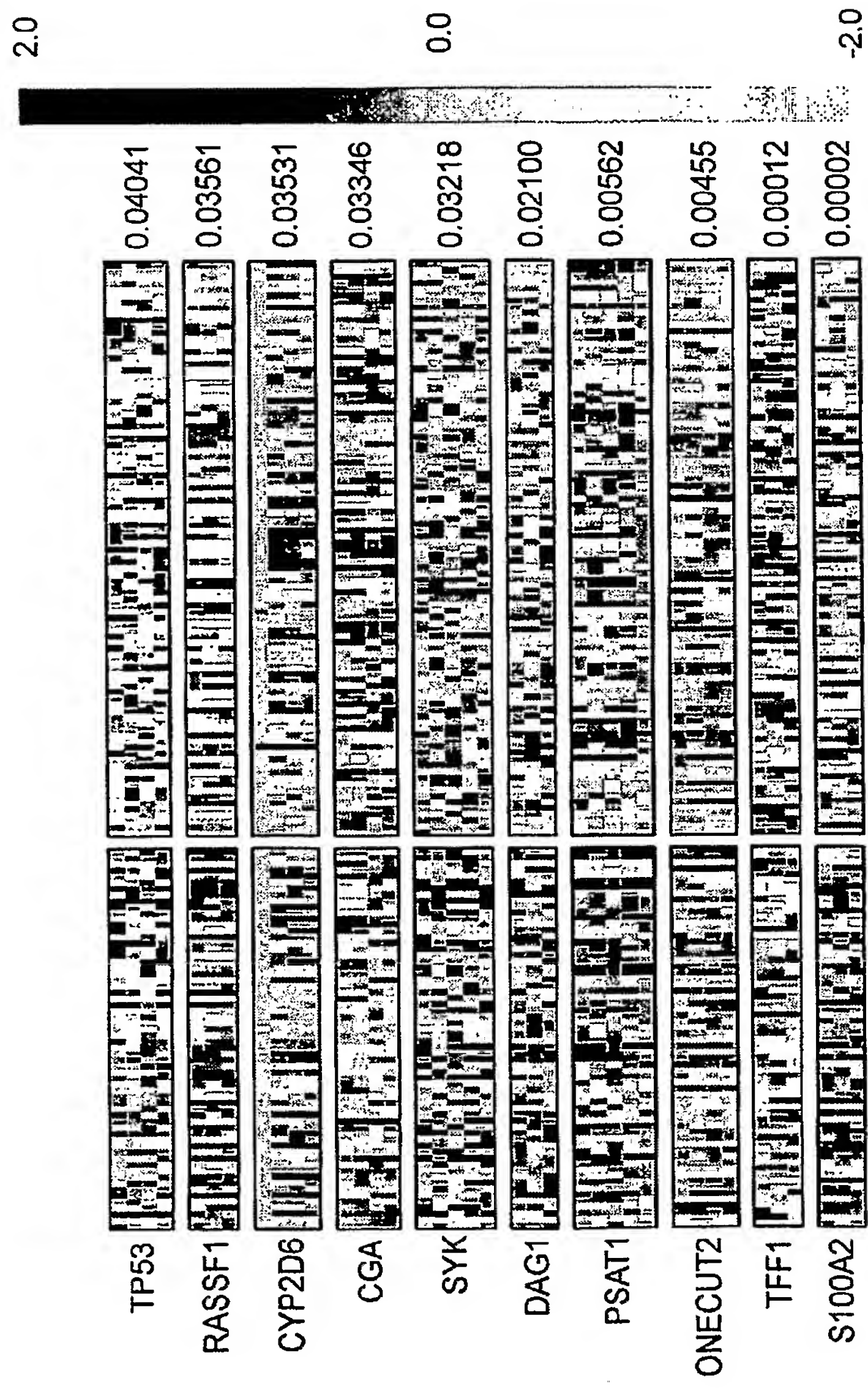


Figure 52 **Stepwise Model (N= 278)**

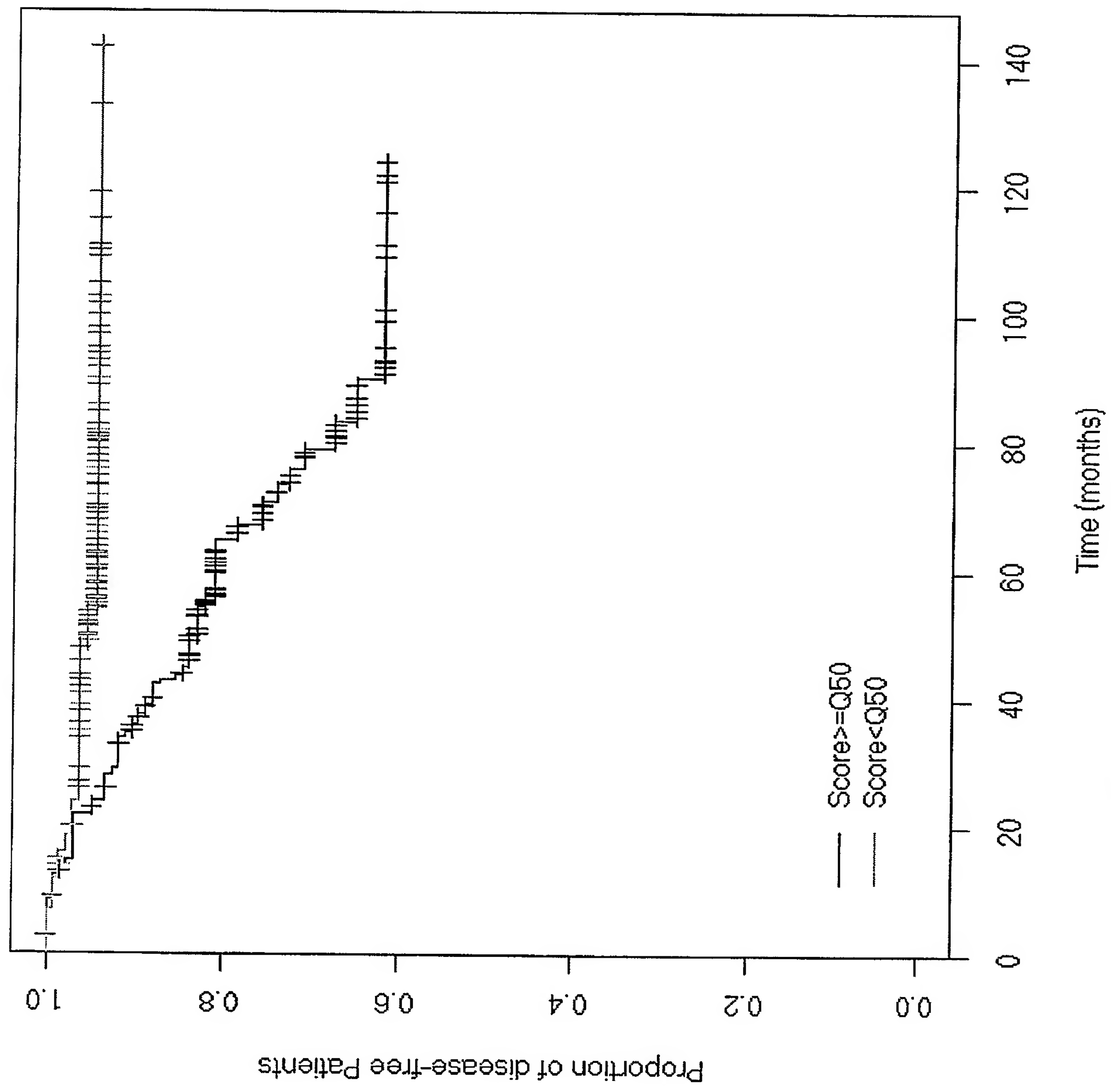


Figure 53 St. Gallen vs. Methylation Marker

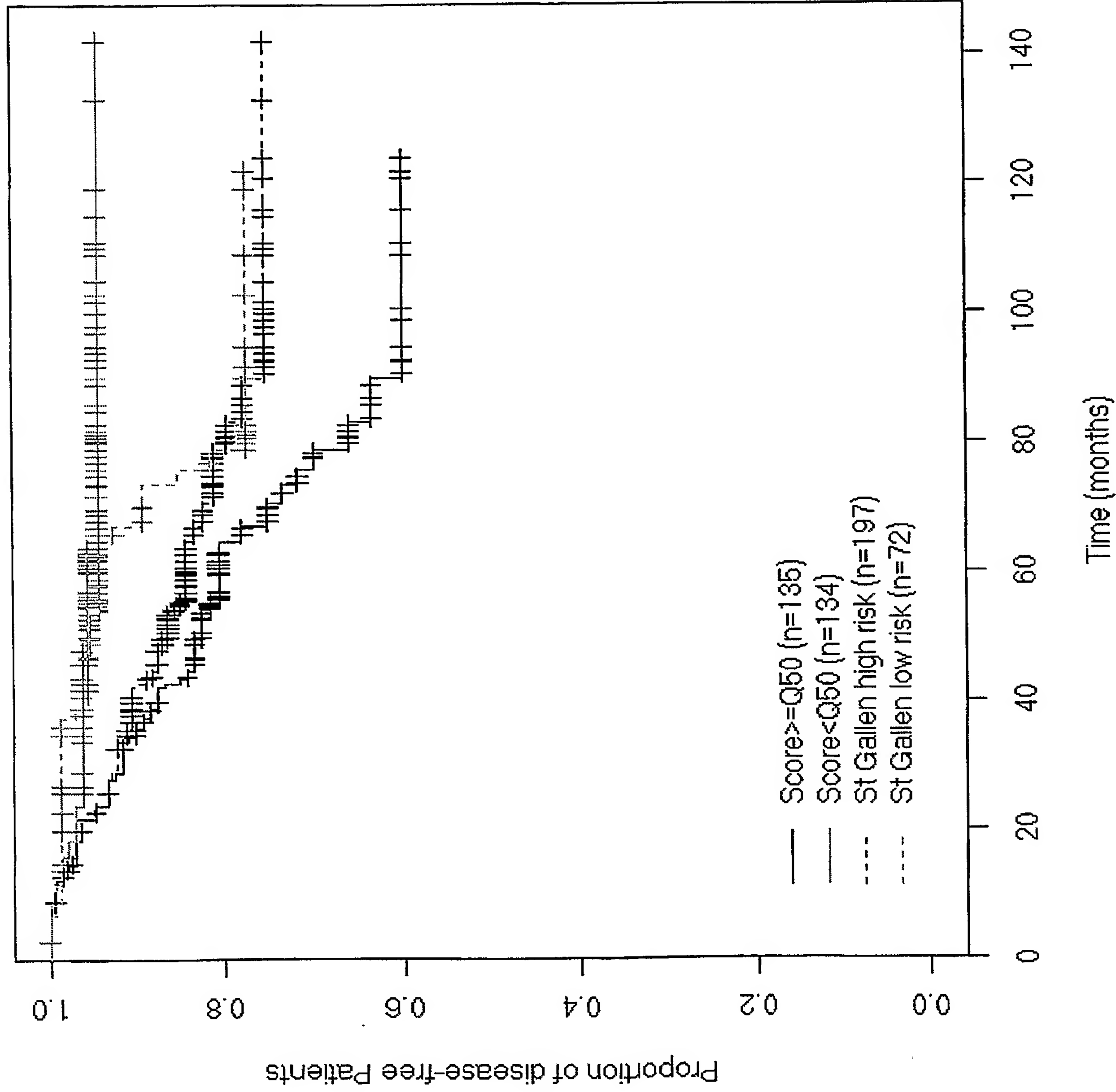
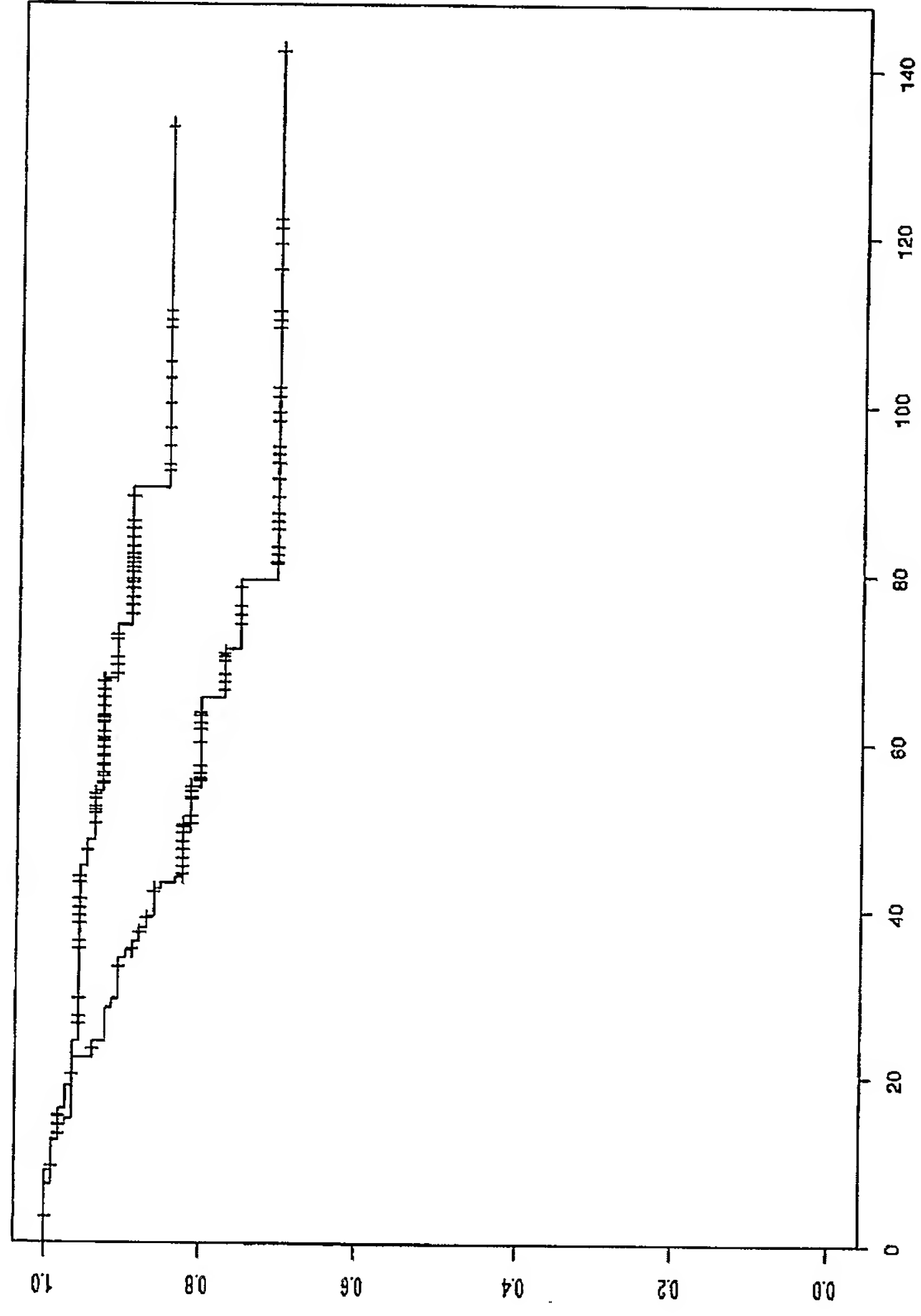


Figure 54



10-02-2004

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<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4491

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 209

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<211> 4256

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 210

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<211> 4256

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4414
<212> DNA
<213> Artificial Sequence

<220>
<223> chemically treated genomic DNA (Homo sapiens)

<400> 212

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<210> 213

<211> 4414

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 213

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<211> 4001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4334

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 216

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4334

<210> 217

<211> 4334

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 217

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<211> 4528

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4374

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4374

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 223

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 10001

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4296

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4296
<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 233

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<211> 4607

<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

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<211> 4453

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 241

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<223> chemically treated genomic DNA (Homo-sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4216

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 244

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<211> 4216

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<223> chemically treated genomic DNA (Homo sapiens)

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<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4448

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<210> 249

<211> 4448

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 257

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<211> 4418

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 258

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<210> 259

<211> 4418

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 259

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<210> 260

<211> 4398

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 260

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<210> 261

<211> 4398

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 261

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<211> 4471

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4492

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 266

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<210> 271

<211> 4001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 271

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<211> 4322

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 272

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4322

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<223> chemically treated genomic DNA (Homo sapiens)

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- <220>
- <223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 277

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 279

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4489

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 286

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4441

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 6435

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

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<210> 315

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<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4337

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 325

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<211> 4388

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<210> 327

<211> 4388

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4256

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4256

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4414

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 337

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4616

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<210> 345

<211> 4374

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 345

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<210> 346

<211> 2534

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 346

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<210> 347

<211> 2534

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 347

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<210> 348

<211> 10001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4449

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 350

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<210> 351

<211> 4449

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 351

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<211> 4296

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<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4296

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<220>
 <223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 355

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<211> 4607

<212> DNA

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<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<210> 364

<211> 3501

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 364

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<210> 365

<211> 3501

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 365

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 aaaataatgt tttattaat tttttttt aggaagata aagatttaa gtatatatt 300
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<210> 366

<211> 4216

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 366

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<210> 367

<211> 4216

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 367

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<210> 372

<211> 4408

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 372

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<210> 373

<211> 4408

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 373

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<211> 4435

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

<400> 374

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4001

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<213> Artificial Sequence

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<400> 378

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<213> Artificial Sequence

<220>
<223> chemically treated genomic DNA (Homo sapiens)

<400> 379

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<211> 4418

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4418

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4479

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 387

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<211> 4492

<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4448

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<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<210> 401

<211> 4001

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 401

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<211> 4341

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<210> 403

<211> 4341

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4433

<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4494

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4489

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 408

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<211> 4489

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

<400> 409

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<211> 4395

<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4499

<212> DNA

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4343

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4343

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4476

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<223> chemically treated genomic DNA (Homo sapiens)

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<211> 4476

<212> DNA

<213> Artificial Sequence

<220>

<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

<213> Artificial Sequence

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<212> DNA

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<211> 4258

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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<223> chemically treated genomic DNA (Homo sapiens)

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17

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16

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agagtttggt ttttgta 18

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attcgtattt gcgggta 18

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<210> 1086

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<400> 1086

aattcgatcg cgggta

16

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<400> 1088

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25

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<400> 1089

aaccaataaa acctactcct cccttaa

27

<210> 1090
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<400> 1090

accaccaccc aacacacaat aacaaacaca 30

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gtaggggagg gaagtagatg tt 22

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agttggagtt gggagagtga aaggaga

27

